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(21) International Application Number: PCT/CZ00/00002 (22) International Filing Date: 25 January 2000 (25.01.00) (30) Priority Data: PV 273-99 26 January 1999 (26.01.99) CZ (71) Applicants (for all designated States except US): ÚSTAV EXPERIMENTÁLNÍ BOTANIKY AV ČR [CZ/CZ]; Rozvojová 135, 160 00 Praha 6 (CZ). LACHEMA, A.S. [CZ/CZ]; Karásek 1, 621 33 Brno (CZ). (72) Inventors; and (75) Inventors/Applicants (for US only): HANUŠ, Jan [CZ/CZ]; Pražského 604, 152 00 Praha 5 (CZ). KRYŠTOF, Vladimír [CZ/CZ]; Křížkova 10, 702 00 Ostrava (CZ). HAJDÚCH, Marián [CZ/CZ]; I.P. Pavlova 18, 775 20 Olomouc (CZ). VESELÝ, Jaroslav [CZ/CZ]; U sokolovny 10, 702 00 Ostrava (CZ). STRNAD, Miroslav [CZ/CZ]; Západní 25, 779 00 Olomouc (CZ). (74) Agent: KUBÁT, Jan; Traplová Hakr Kubát, Law and Patent Offices, Prístavní 24, P.O. Box 38, 170 00 Praha 7 (CZ).		(81) Designated States: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i> <i>Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>
(54) Title: SUBSTITUTED NITROGEN HETEROCYCLIC DERIVATIVES AND PHARMACEUTICAL USE THEREOF (57) Abstract This invention relates to new substituted nitrogen heterocyclic derivatives having cytostatic, anticancer, antimitotic, antineurogenerative, immunosuppressive and antimicrobial effects. This invention also relates to the methods for preparation of these derivatives, the use of the compounds as drugs, the pharmaceutical composition and combined pharmaceutical application form comprising an effective dose of a claimed compound and the use of these derivatives for drug production.		

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Substituted nitrogen heterocyclic derivatives and pharmaceutical use thereof

Field of the invention

This invention relates to new purine and pyrazolopyrimidine derivatives and their
5 deaza analogues and to their use in suitable utilities, especially diagnostic and
therapeutic methods.

It relates, in particular, to purine derivatives and their inhibitory effect with
respect to cyclin-dependent kinase proteins, abbreviated cdks and also with an inhibitory
effect with respect to viruses and immunostimulation. Purine analogues as cdk inhibitors
10 are disclosed for example in WO 97/16452, WO 98/05335 and WO/9720842. The
teaching of these patents includes 2,6,9-trisubstituted and less substituted purine
derivatives only.

Description of related art

15 Tetrasubstituted purines are disclosed in WO 98/01448 in which substituents are
short hydrocarbonyl chains, usually represented by hydrogen. Substituents at C6
represents hydrogen or amine optionally substituted by one or two hydrocarbon groups;
substituents at C8 are hydroxy, mercapto, acyloxy or oxycarbonyl substituted by
aliphatic alkyl only. Nucleotide analogues containing phosphonate groups are disclosed
20 for example in U.S. patents 4,659,825; 4,724,233; 5,124,051; 5,302,585; 5,208,221;
5,352,786; 5,356,886; 5,142,051; in EP publication numbers 269,947; 481,214; 630,381;
369,409; 454,427; 618,214; 398,231; 454,427; 468,119; 481,119; 481,214; 434,450 and
in WO 95/07920; WWO 094/03467, WO96/33200 and WO94/03467. Typical purine
base is adenine, 2,6-diaminopurine and guanine. The purine bases may include the deaza
25 analogues thereof. 6,9-Substituted and 2,6,9-trisubstituted purines and related analogues
are disclosed in WO 96/33200. 2,8,9-, 6,8,9-Trisubstituted and 2,6,8,9-tetrasubstituted
purines and tri- and tetrasubstituted pyrazolopyrimidines and their deaza analogues have
not yet been described.

Summary of the invention

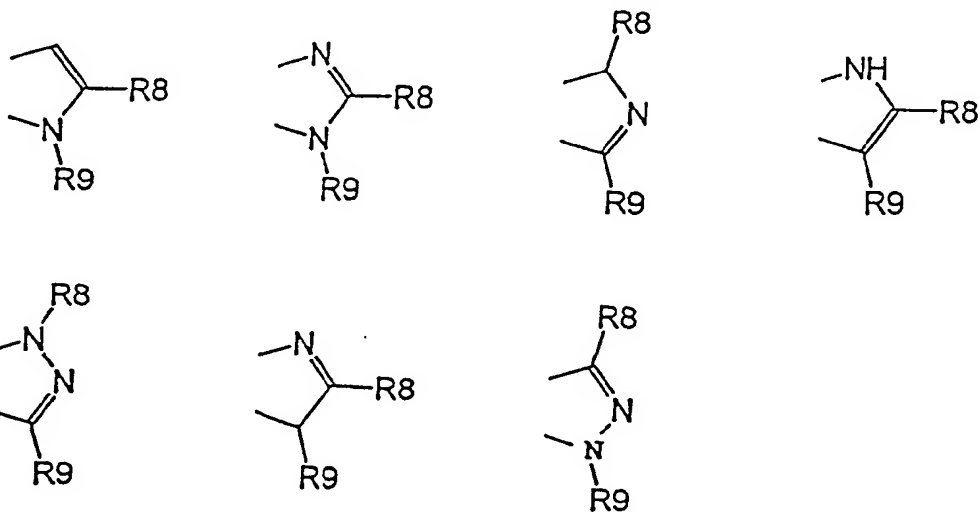
It is an object of this invention to provide anticancer, antiviral, neurodepressive and immunosuppressive compounds having improved selectivity and efficiency index, i.e. that are less toxic yet more efficacious than analogues known heretofore.

The invention concerns substituted nitrogen heterocyclic derivatives of the formula I



wherein,

A is a divalent group selected from the ensemble consisting of



25 Z is N or CH, provided that at least one Z being N;

R2 and R6 are independent of one another, represent H, halogen, alkyl, substituted alkyl, cycloalkyl, substituted cycloalkyl, cycloalkyl alkyl, arylalkyl, heteroalkyl, heteroarylalkyl, cycloheteroalkyl alkyl or R6'-X wherein

X is an -NH-, -N(C₁-C₆-alkyl)-, -O- or -S- moiety;

30 R6' is H, alkyl, substituted alkyl, acyl, amido, sulfo, cycloalkyl, substituted cycloalkyl, aryl, substituted aryl, heterocycle, heteroaryl, substituted heteroaryl, arylalkyl,

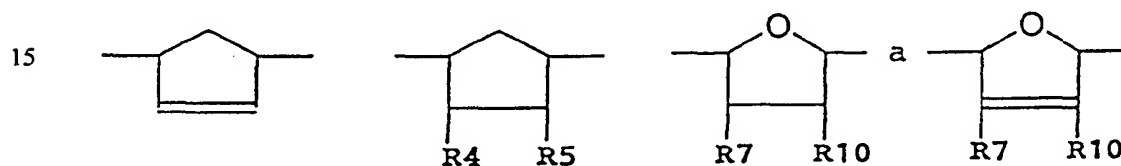
cycloheteroalkyl, substituted cycloheteroalkyl, heteroarylalkyl, heteroalkyl, cycloalkyl alkyl and cycloheteroalkyl alkyl;

R8 is halogen, hydroxyl, amino, carboxyl, cyano, nitro, amido, sulfo, sulfamido, carbamino, alkyl, substituted alkyl, cycloalkyl, substituted cycloalkyl, arylalkyl, heteroalkyl, heteroarylalkyl, cycloalkyl alkyl, cycloheteroalkyl alkyl or **R8'-X** wherein **X** is -NH-, -N(alkyl)-, -O- or -S- moiety and

R8' is according to any one of the substituents defined above for **R2'** or **R6'**.

R9 is alkyl, substituted alkyl, acyl, carboxyl, amido, sulfamino, cycloalkyl, substituted cycloalkyl, cycloalkyl alkyl, heteroalkylcycloalkyl alkyl, heterocycloalkyl, substituted heterocycloalkyl, aryl, substituted aryl, heterocycle, heteroaryl, substituted heteroaryl, arylalkyl, heteroarylalkyl, heteroalkyl or -**B-R9'**- wherein

B is -CH₂-, -(CH₂)₂-, -CH(CH₃)CH₂-, -CH(CH₂F)CH₂-, -CH(CH₂OH)CH₂-, or the groups of the following structure,



wherein the left hand bond is linked to nitrogen of 5-membered ring of compounds of the formula I;

R4 and **R5**, that are independent of one another, represent hydrogen, hydroxyl, halogen, amino, acyloxy substituent having 1-5 carbon atoms, alkoxy substituent having 1-5 carbon atoms, alkylmercapto substituent having 1-5 carbon atoms, alkylamino substituent having 1-5 carbon atoms and dialkylamino in which each alkyl substituent has 1-5 carbon atoms;

R7 and **R10**, that are independent of one another, represent H or alkyl substituent having 1-10 carbon atoms;

R9 is -(CH₂)_n-**R9'**, wherein n = 1-2 and the

R9' is -X(CH₂)_mY wherein

X is -O-, -S-, -NH- or -N(alkyl)- substituent having 1-6 carbon atoms;

m = 1-2;

Y is carboxyl, amido, sulfo, sulfamino, hydroxyl, carboxyl, mercapto, carbylmercapto, amino, alkylamino, carbamino $-\text{PO}(\text{OH})_2$, $-\text{PO}(\text{O}-\text{C}_1-\text{C}_6\text{-alkyl})_2$, $-\text{PO}(\text{NH}-\text{C}_1-\text{C}_6\text{-alkyl})_2$, $\text{PO}(\text{O}-\text{C}_1-\text{C}_6\text{-alkyl})(\text{NH}-\text{C}_1-\text{C}_6\text{-alkyl})$, $-\text{PO}(\text{OH})(\text{O}-\text{C}_1-\text{C}_6\text{-alkyl})$;

5 $-\text{PO}(\text{OH})(\text{NH}-\text{C}_1-\text{C}_6\text{-alkyl})$ or

$-(\text{CH}_2\text{CHD})-\text{R}^9$, wherein

R^9 , X, m and Y are as defined above and

D is alkyl, substituted alkyl, $-\text{PO}(\text{OH})_2$, $-\text{PO}(\text{OH})(\text{O}-\text{C}_1-\text{C}_6\text{-alkyl})$, $-\text{PO}(\text{OH})(\text{NH}-\text{C}_1-\text{C}_6\text{-alkyl})$.

10

The above not yet defined generic group having meanings as introduced in the following legend:

“halogen“

refers to fluorine, bromine, chlorine, and iodine atoms;

15 “alkyl“

refers to

- a branched or unbranched alkyl group having 1-6 carbon atoms,
- a branched or unbranched alkenyl group having 1-6 carbon atoms,
- a branched or unbranched alkynyl group having 1-6 carbon atoms;

20 “substituted alkyl“

refers to a branched or unbranched alkyl, alkenyl or alkynyl group having 1-6 carbon atoms and having substituted by one or more substituents selected from the group consisting of hydroxyl, mercapto, carbylmercapto, halogen, carbyloxy, amino, amido, carboxyl, cycloalkyl, sulfo or acyl. These substituent generic groups having the meanings being identical with the definitions of the corresponding groups as defined in this legend;

25 “carbyloxy“

denotes the group $-\text{OR}_a$, where R_a is alkyl, substituted alkyl, aryl, substituted aryl, arylalkyl, substituted arylalkyl, cycloalkyl, substituted cycloalkyl, heterocycloalkyl or substituted heterocycloalkyl whereas these generic groups have meanings which are identical with definitions of the corresponding groups as defined in this legend;

30 “carbylmercapto“

denotes the group $-SR_b$, where R_b is alkyl, substituted alkyl, aryl, substituted aryl, arylalkyl, substituted arylalkyl, cycloalkyl, substituted cycloalkyl, heterocycloalkyl or substituted heterocycloalkyl whereas these generic groups have meanings which are identical with definitions of the corresponding groups as defined in this legend;

5 "sulfo"

denotes the group $-SO_3R_c$, where R_c is

- hydrogen,
 - a branched or unbranched alkyl group having 1-6 carbon atoms,
 - a branched or unbranched alkenyl group having 1-6 carbon atoms,
 - 10 • a branched or unbranched alkynyl group having 1-6 carbon atoms,
- a branched or unbranched alkyl, alkenyl or alkynyl group having 1-6 carbon atoms and being substituted by one or more substituents selected from the group consisting of hydroxyl, mercapto, carbylmercapto, halogen, carbyloxy, amino, amido, carboxyl, cycloalkyl, sulfo or acyl, whereas these generic groups have meanings which are
- 15 identical with the definitions of the corresponding groups as defined in this legend;

"sulfamino"

denotes the group $-NHSO_3R_d$, wherein R_d is

- hydrogen,
 - a branched or unbranched alkyl group having 1-6 carbon atoms,
 - 20 • a branched or unbranched alkenyl group having 1-6 carbon atoms,
 - a branched or unbranched alkynyl group having 1-6 carbon atoms,
- a branched or unbranched alkyl, alkenyl or alkynyl group having 1-6 carbon atoms and being substituted by one or more substituents selected from the group consisting of hydroxyl, mercapto, carbylmercapto, halogen, carbyloxy, amino, amido, carboxyl,
- 25 cycloalkyl, sulfo or acyl, whereas these generic substituents have meanings which are identical with definitions of the corresponding groups as defined in this legend;

"acyl"

- denotes the group $-C(O)R_e$, where R_e is hydrogen, alkyl, substituted alkyl, aryl, substituted aryl, arylalkyl, substituted arylalkyl, cycloalkyl, substituted cycloalkyl
- 30 whereas these generic groups have meanings which are identical with definitions of the corresponding groups as defined in this legend;

"aryloxy"

denotes the group -OAr, where Ar is an aryl, substituted aryl, heteroaryl or substituted heteroaryl whereas these generic groups have meanings which are identical with definitions of the corresponding groups as defined in this legend;

“alkylamino“

- 5 denotes the group $-NR_fR_g$, where R_f and R_g , that are independent of one another, represent hydrogen, alkyl, substituted alkyl, aryl, substituted aryl, heteroaryl or substituted heteroaryl, whereas these generic substituents have meanings which are identical with definitions of the corresponding groups defined herein;

- “amido“ denotes the group $-C(O)NR_hR_i'$, where R_h and R_i' may independently be
10 hydrogen, alkyl, substituted alkyl, aryl, substituted aryl, heteroaryl, substituted heteroaryl whereas these generic groups have meanings which are identical with definitions of the corresponding groups as defined in this legend;

“carboxyl“

- denotes the group $-C(O)OR_j$, where R_j is hydrogen, alkyl, substituted alkyl, aryl,
15 substituted aryl, heteroaryl or substituted heteroaryl, whereas these generic substituents have meanings which are identical with definitions of the corresponding groups defined herein;

- “carbamino“ denotes the group $-NHCOR_k$, where R_k may be hydrogen, alkyl, substituted alkyl, heterocycle, aryl, substituted aryl, heteroaryl and substituted heteroaryl whereas
20 these generic groups have meanings which are identical with definitions of the corresponding groups as defined in this legend;

“aryl“

- refers to an aromatic carbocyclic group having from 1 to 18 carbon atoms and being composed of at least one aromatic or multiple condensed rings in which at least one of
25 which being aromatic;

“substituted aryl“

- refers to an aromatic carbocyclic group having from 1 to 18 carbon atoms and being composed of at least one aromatic ring or of multiple condensed rings at least one of which being aromatic. The ring(s) are optionally substituted with one or more
30 substituents selected from the group consisting of halogen, alkyl, hydroxyl, carbylmercapto, alkylamino, carbyloxy, amino, amido, carboxyl, nitro, mercapto or

sulfo, whereas these generic substituent group have meanings which are identical with definitions of the corresponding groups as defined in this legend;

“heterocycle“

refers to a heterocyclic group having from 4 to 9 carbon atoms and at least one
5 heteroatom selected from the group consisting of N, O or S;

“heteroaryl“

refers to a heterocyclic group having from 4 to 9 carbon atoms and at least one
heteroatom selected from the group consisting of N, O or S with at least one ring of this
group being aromatic;

10 “substituted heteroaryl“

refers to a heterocyclic group having from 4 to 9 carbon atoms and at least one
heteroatom selected from the group consisting of N, O or S with at least one ring of this
group being aromatic and this group being substituted with one or more substituents
selected from the group consisting of halogen, alkyl, carbyloxy, carbylmercapto,
15 alkylamino, amido, carboxyl, hydroxyl, nitro, mercapto or sulfo, whereas these generic
substituent group have meanings which are identical with definitions of the
corresponding groups as defined in this legend;

“arylalkyl“

denotes the group $-R_1-Ar$ where R_1 is

- 20
- a branched or unbranched alkyl group having 1-6 carbon atoms,
 - a branched or unbranched alkenyl group having 1-6 carbon atoms,
 - a branched or unbranched alkynyl group having 1-6 carbon atoms,

and Ar is

an aromatic carbocyclic group having from 1 to 18 carbon atoms and being composed of
25 at least one aromatic ring or of multiple condensed rings at least one of which being
aromatic and the group being optionally substituted with one or more substituents
selected from the group consisting of halogen, alkyl, hydroxyl, carbylmercapto,
alkylamino, carbyloxy, amino, amido, carboxyl, nitro, mercapto or sulfo, whereas these
generic substituent group have meanings which are identical with definitions of the
30 corresponding groups as defined in this legend;

“heteroalkyl“

denotes the group $-R_m-L$ where L is

- a branched or unbranched alkyl group having 1-6 carbon atoms,
- a branched or unbranched alkenyl group having 1-6 carbon atoms,
- a branched or unbranched alkynyl group having 1-6 carbon atoms,
- a branched or unbranched alkyl, alkenyl or alkynyl group having 1-6 carbon atoms and being substituted by one or more substituents selected from the group consisting of hydroxyl, mercapto, carbylmercapto, halogen, carbyloxy, amino, amido, carboxyl, cycloalkyl, sulfo or acyl, whereas these generic substituent group have meanings which are identical with definitions of the corresponding groups as defined in this legend

10 and L is

a heterocyclic group having from 4 to 9 carbon atoms and at least one heteroatom selected from the group consisting of N, O or S and the group being unsubstituted or substituted with one or more substituents selected from the group consisting of halogen, alkyl, alkoxy, alkylmercapto, alkylamino, amido, carboxyl, hydroxy, nitro, mercapto or sulfo, whereas these generic substituents have meanings which are identical with definitions of the corresponding groups as defined in this legend;

“heteroarylalkyl”

denotes the group -R_n-G- where R_n is

- a branched or unbranched alkyl group having 1-6 carbon atoms,
- a branched or unbranched alkenyl group having 1-6 carbon atoms,
- a branched or unbranched alkynyl group having 1-6 carbon atoms,
- a branched or unbranched alkyl, alkenyl or alkynyl group having 1-6 carbon atoms and being substituted by one or more substituents selected from the group consisting of hydroxyl, mercapto, carbylmercapto, halogen, carbyloxy, amino, amido, carboxyl, cycloalkyl, sulfo or acyl, whereas these generic substituent group have meanings which are identical with definitions of the corresponding groups as defined in this legend

and G is

a heterocyclic group having from 4 to 9 carbon atoms and at least one heteroatom selected from the group consisting of N, O or S with at least one ring of which being aromatic and the group being optionally substituted with one or more substituents selected from the group consisting of halogen, alkyl, carbyloxy, carbylmercapto,

alkylamino, amido, carboxyl, hydroxyl, nitro, mercapto or sulfo, whereas these generic substituent group have meanings which are identical with definitions of the corresponding groups as defined in this legend;

“cycloalkyl“

5 refers to a monocyclic or polycyclic alkyl group containing 3 to 15 carbon atoms;

“substituted cycloalkyl“

refers to a monocyclic or polycyclic alkyl group containing 3 to 15 carbon atoms and being substituted by one or more substituents selected from the group consisting of halogen, alkyl, substituted alkyl, carbyloxy, carbylmercapto, aryl, nitro, mercapto or
10 sulfo, whereas these generic substituent groups have meanings which are identical with definitions of the corresponding groups as defined in this legend;

“heterocycloalkyl“

refers to a monocyclic or polycyclic alkyl group containing 3 to 15 carbon atoms which at least one ring carbon atom of its cyclic structure being replaced with a heteroatom
15 selected from the group consisting of N, O, S or P;

“substituted heterocycloalkyl“

refers to a monocyclic or polycyclic alkyl group containing 3 to 15 carbon atoms which at least one ring carbon atom of its cyclic structure being replaced with a heteroatom selected from the group consisting of N, O, S or P and the group is containing one or
20 more substituents selected from the group consisting of halogen, alkyl, substituted alkyl, carbyloxy, carbylmercapto, aryl, nitro, mercapto or sulfo, whereas these generic substituent group have meanings which are identical with definitions of the corresponding groups as defined in this legend;

“cycloalkyl alkyl“

25 denotes the group $-R_o-J$ where R_o is

- a branched or unbranched alkyl group having 1-6 carbon atoms,
- a branched or unbranched alkenyl group having 1-6 carbon atoms,
- a branched or unbranched alkynyl group having 1-6 carbon atoms,
- a branched or unbranched alkyl, alkenyl or alkynyl group having 1-6 carbon atoms
30 and being substituted by one or more substituents selected from the group consisting of hydroxyl, mercapto, carbylmercapto, halogen, carbyloxy, amino, amido, carboxyl, cycloalkyl, sulfo or acyl, whereas these generic substituent group have meanings

which are identical with definitions of the corresponding groups as defined in this legend

and J is

- a monocyclic or polycyclic alkyl group containing 3 to 15 carbon atoms;
- a monocyclic or polycyclic alkyl group containing 3 to 15 carbon atoms which contains one or more substituents selected from the group consisting of halogen; alkyl, substituted alkyl, carbyloxy, carbylmercapto, aryl, nitro, mercapto or sulfo, whereas these generic substituent groups have meanings which are identical with definitions of the corresponding groups as defined in this legend and

“heterocycloalkylalkyl”

denotes the group $-R_pV$ where R_p is

- a branched or unbranched alkyl group having 1-6 carbon atoms,
- a branched or unbranched alkenyl group having 1-6 carbon atoms,
- a branched or unbranched alkynyl group having 1-6 carbon atoms,
- a branched or unbranched alkyl, alkenyl or alkynyl group having 1-6 carbon atoms and being substituted by one or more substituents selected from the group consisting of hydroxyl, mercapto, carbylmercapto, halogen, carbyloxy, amino, amido, carboxyl, cycloalkyl, sulfo or acyl, whereas these generic substituent groups have meanings which are identical with definitions of the corresponding groups as defined in this

legend

and V is

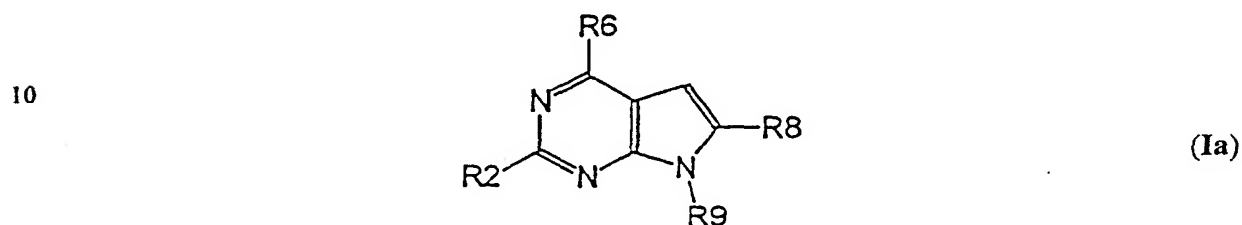
- a monocyclic or polycyclic alkyl group containing 3 to 15 carbon atoms with at least one being replaced with a heteroatom selected from the group consisting of N, O, S or P;
- a monocyclic or polycyclic alkyl group containing 3 to 15 carbon atoms with at least one being replaced with a heteroatom selected from the group consisting of N, O, S or P and the group contains one or more substituents selected from the group consisting of halogen, alkyl, substituted alkyl, carbyloxy, carbylmercapto, aryl, nitro, mercapto or sulfo, whereas these generic substituent groups have meanings which are identical with definitions of the corresponding groups as defined in this legend

and the pharmaceutically acceptable acid salts, racemates and optical isomers thereof.

The invention concerns further substituted nitrogen heterocyclic derivatives of the formula **I**, wherein R6 = H and R2, R8 and R9 have above mentioned meanings.

The invention concerns further substituted nitrogen heterocyclic derivatives of the formula **I**, wherein R2 = H and R6, R8 and R9 have above mentioned meanings.

The invention concerns further substituted nitrogen heterocyclic derivatives of the formula **Ia**,

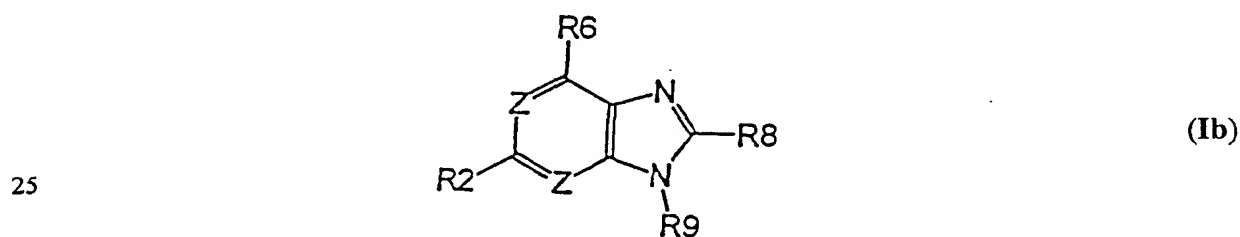


wherein R2, R6, R8 and R9 have above mentioned meanings.

15 The invention concerns further substituted nitrogen heterocyclic derivatives of the formula **Ia**, wherein R6 = H and R2, R8 and R9 have above mentioned meanings.

The invention concerns further substituted nitrogen heterocyclic derivatives of the formula **Ia**, wherein R2 = H and R6, R8 and R9 have above mentioned meanings.

20 The invention concerns further substituted nitrogen heterocyclic derivatives of the formula **Ib**,



wherein R2, R6, R8 and R9 have above mentioned meanings.

30 The invention concerns further substituted nitrogen heterocyclic derivatives of the formula **Ib**, wherein R6 = H and R2, R8 and R9 have above mentioned meanings.

The invention concerns further substituted nitrogen heterocyclic derivatives of the formula **Ib**, wherein R2 = H and R6, R8 and R9 have above mentioned meanings.

The invention concerns further substituted nitrogen heterocyclic derivatives of the formula **Ic**,

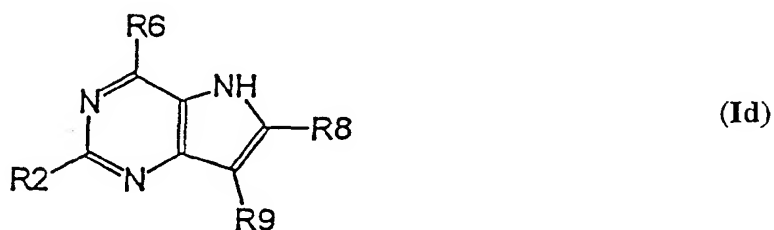


wherein R2, R6, R8 and R9 have above mentioned meanings.

10 The invention concerns further substituted nitrogen heterocyclic derivatives of the formula **Ic**, wherein R6 = H and R2, R8 and R9 have above mentioned meanings.

The invention concerns further substituted nitrogen heterocyclic derivatives of the formula **Ic**, wherein R2 = H and R6, R8 and R9 have above mentioned meanings.

15 The invention concerns further substituted nitrogen heterocyclic derivatives of the formula **Id**

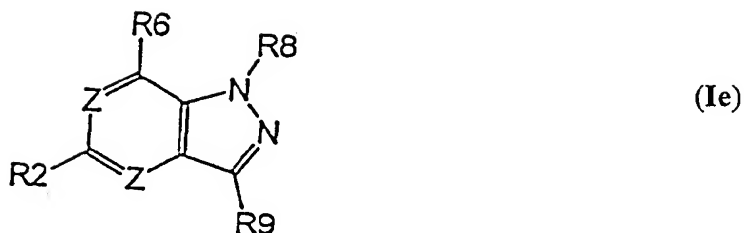


wherein R2, R6, R8 and R9 have above mentioned meanings.

25 The invention concerns further substituted nitrogen heterocyclic derivatives of the formula **Id**, wherein R6 = H and R2, R8 and R9 have above mentioned meanings.

The invention concerns further substituted nitrogen heterocyclic derivatives of the formula **Id**, wherein R2 = H and R6, R8 and R9 have above mentioned meanings.

30 The invention concerns further substituted nitrogen heterocyclic derivatives of the formula **Ie**,

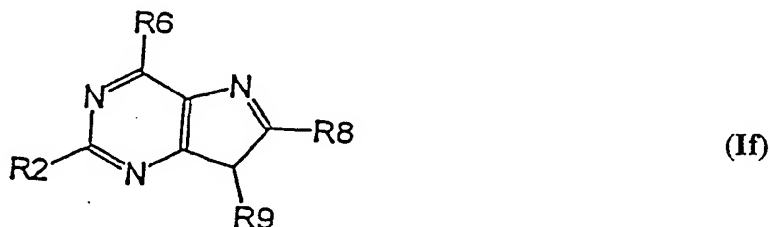


wherein R2, R6, R8 and R9 have above mentioned meanings.

The invention concerns further substituted nitrogen heterocyclic derivatives of the formula **Ie**, wherein R6 = H and R2, R8 and R9 have above mentioned meanings.

10 The invention concerns further substituted nitrogen heterocyclic derivatives of the formula **Ie**, wherein R2 = H and R6, R8 and R9 have above mentioned meanings.

The invention concerns further substituted nitrogen heterocyclic derivatives of the formula **If**,



20 wherein R2, R6, R8 and R9 have above mentioned meanings.

The invention concerns further substituted nitrogen heterocyclic derivatives of the formula **If**, wherein R6 = H and R2, R8 and R9 have above mentioned meanings.

The invention concerns further substituted nitrogen heterocyclic derivatives of the formula **If**, wherein R2 = H and R6, R8 and R9 have above mentioned meanings.

25 The invention concerns further substituted nitrogen heterocyclic derivatives of the formula **Ig**,



wherein R2, R6, R8 and R9 have above mentioned meanings.

The invention concerns further substituted nitrogen heterocyclic derivatives of the formula **Ig**, wherein R6 = H and R2, R8 and R9 have above mentioned meanings.

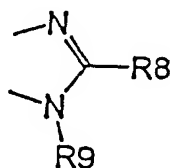
The invention concerns further substituted nitrogen heterocyclic derivatives of the formula **Ig**, wherein R2 = H and R6, R8 and R9 have above mentioned meanings.

The invention concerns further substituted nitrogen heterocyclic derivatives of the formula **I** selected from the group consisting of 2-(1-hydroxymethylpropylamino)-6-benzylamino-8-chloro (or hydroxy, bromo, fluoro, amino, amido, carboxy, cyano, methylamino, thio, methylthio, ω -hydroxyalkylamino, ω -hydroxyalkyloxy, ω -carboxyalkylamino, ω -aminoalkylamino, ω -fosfonoalkylamino, ω -fosfonoalkyloxy, propinyl)-9-isopropylpurine, 2-(2-aminopropylamino)-6-benzylamino-8-chloro (or hydroxy, bromo, fluoro, amino, amido, carboxy, cyano, methylamino, thio, methylthio, ω -hydroxyalkylamino, ω -hydroxyalkyloxy, ω -carboxyalkylamino, ω -aminoalkylamino, ω -fosfonoalkylamino, ω -fosfonoalkyloxy, propinyl)-9-isopropylpurine, 2-(2-hydroxypropylamino)-6-benzylamino-8-chloro (or hydroxy, bromo, fluoro, amino, amido, carboxy, cyano, methylamino, thio, methylthio, ω -hydroxyalkylamino, ω -hydroxyalkyloxy, ω -carboxyalkylamino, ω -aminoalkylamino, ω -fosfonoalkylamino, ω -fosfonoalkyloxy, propinyl)-9-isopropylpurine, 2-diethylamino-6-(4-methoxybenzylamino)-8-chloro (or hydroxy, bromo, fluoro, amino, amido, carboxy, cyano, methylamino, thio, methylthio, ω -hydroxyalkylamino, ω -hydroxyalkyloxy, ω -carboxyalkylamino, ω -aminoalkylamino, ω -fosfonoalkylamino, ω -fosfonoalkyloxy, propinyl)-9-isopropylpurine, 2-(2-hydroxypropylamino)-6-(3-chloroanilino)-8-chloro (or hydroxy, bromo, amino, C₁-C₆ alkyl, methyl, ethyl, propyl, isopropyl, vinyl, allyl, propargyl)-9-isopropylpurine, 2-(2-hydroxypropylamino)-6-(3-chloro-4-carboxyanilino)-8-chloro (or hydroxy, bromo, fluoro, amino, amido, carboxy, cyano, methylamino, thio, methylthio, ω -hydroxyalkylamino, ω -hydroxyalkyloxy, ω -carboxyalkylamino, ω -aminoalkylamino, ω -fosfonoalkylamino, ω -fosfonoalkyloxy, propinyl)-9-isopropylpurine, 2-(R)-(2-hydroxypyrrolidin-1-yl)-6-benzylamino-8-chloro (or hydroxy, bromo, fluoro, amino, amido, carboxy, cyano, methylamino, thio, methylthio, ω -hydroxyalkylamino, ω -hydroxyalkyloxy, ω -carboxyalkylamino, ω -aminoalkylamino, ω -fosfonoalkylamino, ω -fosfonoalkyloxy, propinyl)-9-isopropylpurine, 2-(R)-(1-isopropyl-2-hydroxyethylamino)-6-(3-chloro-4-carboxyanilino)-8-chloro (or hydroxy, bromo, fluoro, amino, amido,

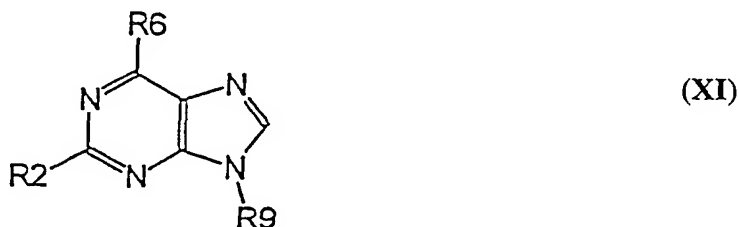
carboxy, cyano, methylamino, thio, methylthio, ω -hydroxyalkylamino, ω -hydroxyalkyloxy, ω -carboxyalkylamino, ω -aminoalkylamino, ω -fosfonoalkylamino, ω -fosfonoalkyloxy, propinyl)-9-isopropylpurine, 2-(*R*)-(1-isopropyl-2-hydroxyethylamino)-6-benzylamino-8-chloro (or hydroxy, bromo, fluoro, amino, amido, carboxy, cyano, methylamino, thio, methylthio, ω -hydroxyalkylamino, ω -hydroxyalkyloxy, ω -carboxyalkylamino, ω -aminoalkylamino, ω -fosfonoalkylamino, ω -fosfonoalkyloxy, propinyl)-9-isopropylpurine, 2-(*R*)-(1-isopropyl-2-hydroxyethylamino)-6-(3-chloroanilino)-8-chloro (or hydroxy, bromo, fluoro, amino, amido, carboxy, cyano, methylamino, thio, methylthio, ω -hydroxyalkylamino, ω -hydroxyalkyloxy, ω -carboxyalkylamino, ω -aminoalkylamino, ω -fosfonoalkylamino, ω -fosfonoalkyloxy, propinyl)-9-isopropylpurine, 2-alkylamino-6-dimethylamino-8-chloro (or hydroxy, bromo, fluoro, amino, amido, carboxy, cyano, methylamino, thio, methylthio, ω -hydroxyalkylamino, ω -hydroxyalkyloxy, ω -carboxyalkylamino, ω -aminoalkylamino, ω -fosfonoalkylamino, ω -fosfonoalkyloxy, propinyl)-9-(*R*)-(2-phosphonomethoxypropyl)purine, 2-alkylamino-6-diethylamino-8-chloro (or hydroxy, bromo, fluoro, amino, amido, carboxy, cyano, methylamino, thio, methylthio, ω -hydroxyalkylamino, ω -hydroxyalkyloxy, ω -carboxyalkylamino, ω -aminoalkylamino, ω -fosfonoalkylamino, ω -fosfonoalkyloxy, propinyl)-9-(*R*)-(2-phosphonomethoxypropyl)purine, 2-alkylamino-6-butylamino-8-chloro (or hydroxy, bromo, fluoro, amino, amido, carboxy, cyano, methylamino, thio, methylthio, ω -hydroxyalkylamino, ω -hydroxyalkyloxy, ω -carboxyalkylamino, ω -aminoalkylamino, ω -fosfonoalkylamino, ω -fosfonoalkyloxy, propinyl)-9-(*R*)-(2-phosphonomethoxypropyl)purine, 2-alkylamino-6-(2-butylamino)-8-chloro (or hydroxy, bromo, fluoro, amino, amido, carboxy, cyano, methylamino, thio, methylthio, ω -hydroxyalkylamino, ω -hydroxyalkyloxy, ω -carboxyalkylamino, ω -aminoalkylamino, ω -fosfonoalkylamino, ω -fosfonoalkyloxy, propinyl)-9-(*R*)-(2-phosphonomethoxypropyl)purine, 2-alkylamino-6-cyclopropylamino-8-chloro (or hydroxy, bromo, fluoro, amino, amido, carboxy, cyano, methylamino, thio, methylthio, ω -hydroxyalkylamino, ω -hydroxyalkyloxy, ω -carboxyalkylamino, ω -aminoalkylamino, ω -fosfonoalkylamino, ω -fosfonoalkyloxy, propinyl)-9-(*R*)-(2-phosphonomethoxypropyl)purine, 2-amino-6-cyclohexylamino-8-chloro (or hydroxy, bromo, fluoro, amino, amido, carboxy, cyano, methylamino, thio, methylthio, ω -hydroxyalkylamino, ω -hydroxyalkyloxy, ω -carboxyalkylamino, ω -aminoalkylamino, ω -fosfonoalkylamino, ω -fosfonoalkyloxy, propinyl)-9-(*R*)-(2-

phosphonomethoxypropyl)purine, 2-alkylamino-6-(pyrrolidin-1-yl)-8-chloro (or hydroxy, bromo, fluoro, amino, amido, carboxy, cyano, methylamino, thio, methylthio, ω -hydroxyalkylamino, ω -hydroxyalkyloxy, ω -carboxyalkylamino, ω -aminoalkylamino, ω -fosfonoalkylamino, ω -fosfonoalkyloxy, propinyl)-9-(R)-(2-phosphonomethoxypropyl)purine, 2-alkylamino-6-(morpholin-1-yl)-8-chloro (or hydroxy, bromo, fluoro, amino, amido, carboxy, cyano, methylamino, thio, methylthio, ω -hydroxyalkylamino, ω -hydroxyalkyloxy, ω -carboxyalkylamino, ω -aminoalkylamino, ω -fosfonoalkylamino, ω -fosfonoalkyloxy, propinyl)-9-(R)-(2-phosphonomethoxypropyl)purine.

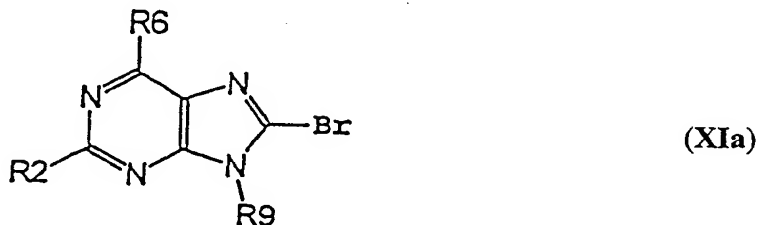
The invention also relates to a method for preparing substituted nitrogen heterocyclic derivatives of formula I, wherein R₂, R₆, R₈ and R₉ have above mentioned meanings, A is a group of formula



and Z is N, characterized in that a trisubstituted derivative of formula XI,

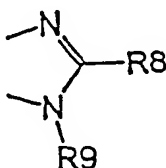


wherein R₂, R₆ and R₉ have above mentioned meanings, is brominated with using a brominating system selected from a group consisting of bromine/acetic acid, bromine/chloroform, bromine/acetate buffer, bromine/water, N-bromosuccinimide/dimethylformamide and bromoacetamide/dimethylformamide to obtain a derivative of formula XIa,

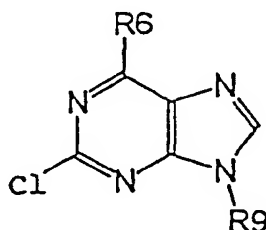


wherein R2, R6 and R9 have above mentioned meanings, and the bromine atom in position 8 of the derivative of formula **XIa** is then optionally subjected to a substitution in order to replace it by another substituent R8, which has above mentioned meanings.

The invention also relates to a method for preparing substituted nitrogen heterocyclic derivatives of formula **I**, wherein R2, R6, R8 and R9 have above mentioned meanings, A is a group of formula

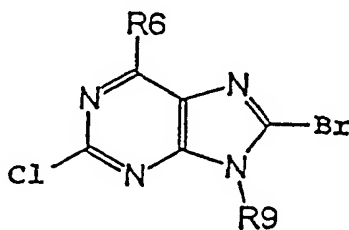


and Z is N, characterized in that a trisubstituted derivative of formula **XIb**,



(**XIb**)

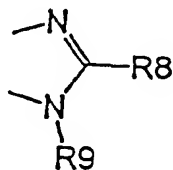
wherein R6 and R9 have above mentioned meanings, is brominated with using a brominating system selected from a group consisting of bromine/acetic acid, bromine/chloroform, bromine/acetate buffer, bromine/water, N-bromosuccinimide/dimethylformamide and bromoacetamide/dimethylformamide to obtain a derivative of formula **XIc**,



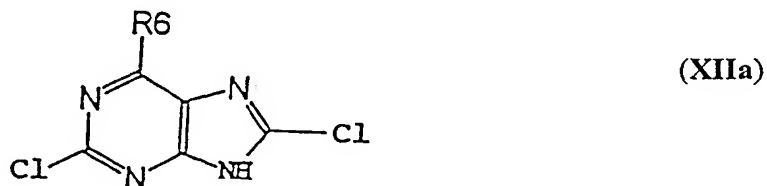
(**XIc**)

wherein R6 and R9 have above mentioned meanings, and the chlorine atom in position 2 and the bromine atom in position 8 of the derivative of formula **XIc** are optionally, either progressively or simultaneously, subjected to a nucleophilic substitution in order to replace them by another substituents R2 and R8, that have above mentioned meanings.

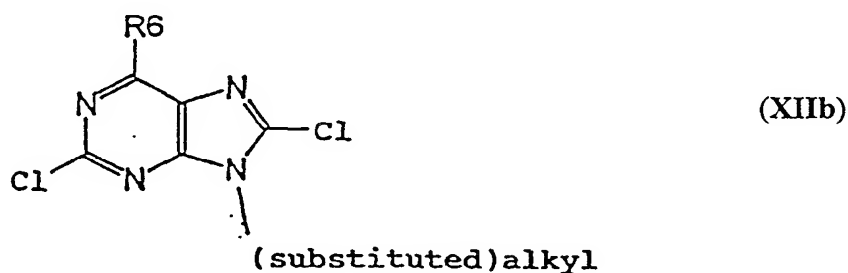
The invention also relates to a method for preparing substituted nitrogen heterocyclic derivatives of formula I, wherein R₂, R₆ and R₈ have above mentioned meanings, R₉ is alkyl as defined above, A is a group of formula



and Z is N, characterized in that, a trisubstituted derivative of formula XIIa,



wherein R₆ is as defined above, is alkylated in position 9 with using an appropriate alkylating agent in a system selected from a group consisting of K₂CO₃/dimethylformamide, Cs₂CO₃/dimethylformamide, t-BuOK/dimethylformamide, t-BuOK/dimethylsulphoxide and NaH/dimethylformamide or under conditions of Mitsunobu reaction to obtain a derivative of formula XIIb

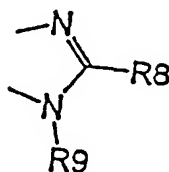


wherein R₆ and alkyl are as defined above, and the chlorine atoms in positions 2 and 8 of the derivative of formula XIIb are optionally, either progressively or simultaneously, subjected to a substitution in order to replace them by another substituents R₂ and R₈, that have above mentioned meanings.

The invention also relates to a method for preparing substituted nitrogen heterocyclic derivatives, wherein R₂, R₆ and R₈ have above mentioned meanings, R₉ is

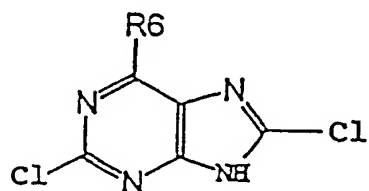
alkyl as defined above, A is a group of formula

5



and Z is N, characterized in that a trisubstituted derivative of formula **XIIa**,

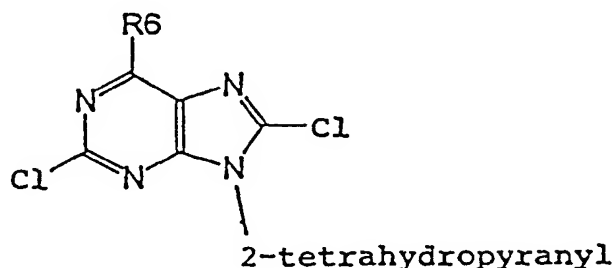
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(XIIa)

15 wherein R6 has above mentioned meanings, is protected by reacting, for example, with 2-dihydropyrane/ H^+ to obtain a derivative of formula **XIIc**,

20



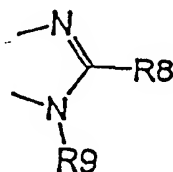
(XIIc)

wherein R6 has above mentioned meanings, and a group R6 and/or the chlorine atom in position 2 and/or the chlorine atom in position 8 are optionally converted to another substituents R2, R6 and R8, that have above mentioned meanings and in such optionally modified product of formula **XIIc**, the 2-tetrahydropyranyl group is split off to obtain a corresponding derivative of formula **XIIc** wherein R9 is hydrogen, and so obtained product is then alkylated in position 9 with using an appropriate alkylating agent in a system selected from a group consisting of K_2CO_3 /dimethylformamide, Cs_2CO_3 /dimethylformamide, t-BuOK/dimethylformamide, t-BuOK/dimethylsulphoxide and NaH/dimethylformamide or under conditions of Mitsunobu reaction to obtain a tetrasubstituted derivative of formula **XIIc**, wherein R2, R6, R8 and R9 have above

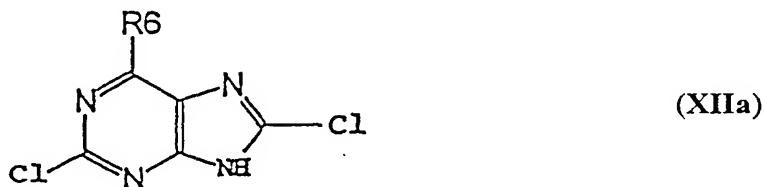
30

mentioned meanings.

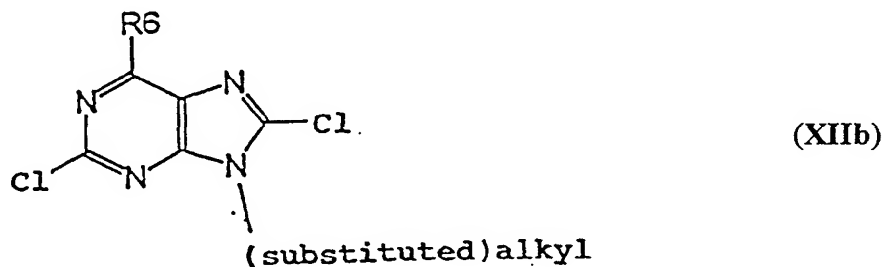
The invention also relates to a method for preparing substituted nitrogen heterocyclic derivatives of formula I, wherein R2, R6 and R8 have above mentioned meanings, R9 is alkyl as defined above, A is a group



and Z is N, characterized in that, 2,6,8-trichloropurine is subjected to a nucleophilic substitution in position 6 in order to replace the chlorine atom in position 6 by another substituent R6 as defined above to obtain a derivative of formula XIIa,



wherein R6 has above mentioned meanings, and so obtained product is then alkylated in position 9 with using an appropriate alkylating agent in a system selected from a group consisting of K_2CO_3 /dimethylformamide, Cs_2CO_3 /dimethylformamide, t-BuOK/dimethylformamide, t-BuOK/dimethylsulphoxide and NaH/dimethylformamide or under conditions of Mitsunobu reaction to obtain a derivative of formula XIIb,



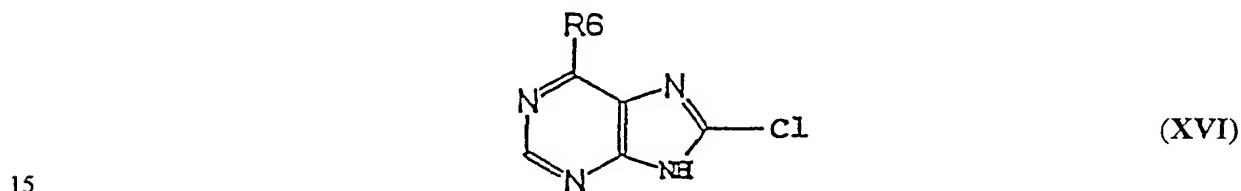
wherein R6 and alkyl have above mentioned meanings, and the chlorine atoms in positions 2 and 8 of the derivative of formula XIIb are optionally, either progressively or simultaneously, subjected to a nucleophilic substitution in order to replace them by another substituents R2 and R8 that have above mentioned meanings.

The invention also relates to a method for preparing substituted nitrogen heterocyclic derivatives of formula I, wherein R2 is hydrogen, R6 and R8 have above mentioned meanings, R9 is alkyl as defined above, A is a group of formula



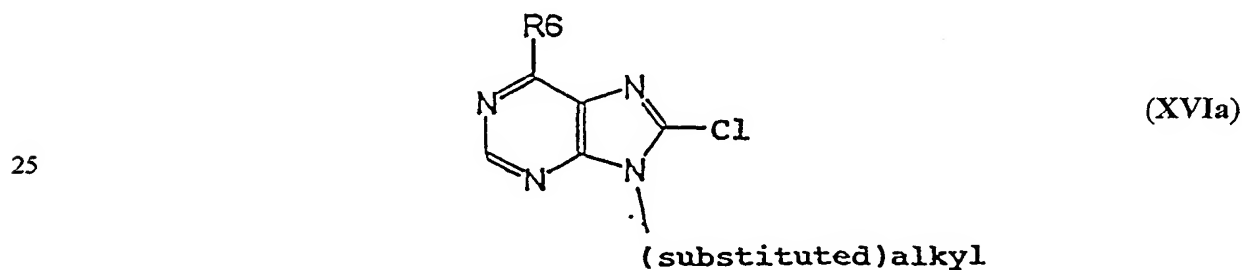
and Z is N, characterized in that 6,8-dichloropurine is subjected to a substitution in position 6 in order to replace the chlorine atom in position 6 by another substituent R6 which has above mentioned meanings to obtain a derivative of formula XVI,

10



wherein R6 has above mentioned meanings, and so obtained derivative is alkylated in position 9 with using an appropriate alkylating agent in a system selected from a group consisting of K₂CO₃/dimethylformamide, Cs₂CO₃/dimethylformamide, t-BuOK/dimethylformamide, t-BuOK/dimethylsulphoxide and NaH/dimethylformamide or under conditions of Mitsunobu reaction to obtain a derivative of formula XVIa,

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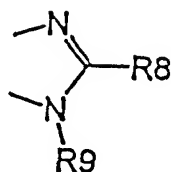


wherein R6 and alkyl have above mentioned meanings, and the chlorine atom in position 8 of the derivative of formula XVIa is optionally subjected to a substitution in order to replace it by another substituent R8 which has above mentioned meanings.

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The invention also relates to a method for preparing substituted nitrogen heterocyclic derivatives of formula I, wherein R2, R6 and R8 have above mentioned

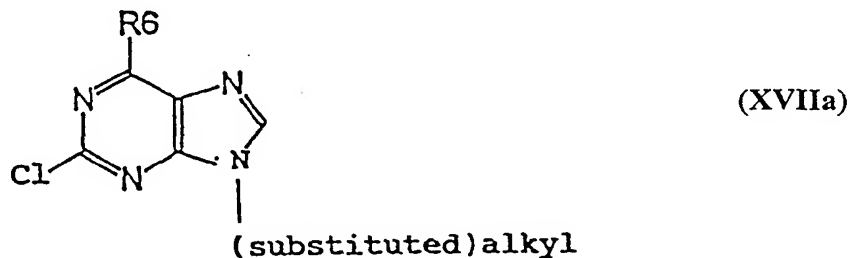
meanings, R₉ is alkyl as defined above, A is a group of formula



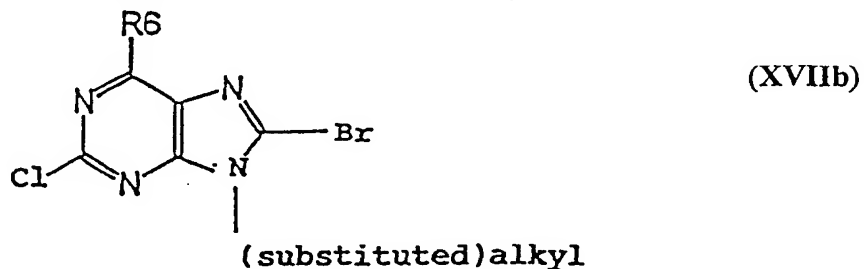
and Z is N, characterized in that 2,6-dichloropurine is subjected to a nucleophilic substitution in position 6 in order to replace the chlorine atom in position 6 by another substituent R₆ as defined above, and so obtained derivative of formula XVII,



wherein R₆ is as defined above, is alkylated in position 9 with using either methyl acrylate or acrylonitrile or oxirane to obtain a derivative of formula XVIIa

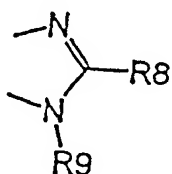


wherein R₆ and alkyl have above mentioned meanings, which product is then brominated in position 8 with using a brominating system selected from a group consisting of bromine/acetic acid, bromine/chloroform, bromine/acetate buffer, bromine/water, N-bromosuccinimide/dimethylformamide and bromoacetamide/dimethylformamide to obtain a derivative of formula XVIIb,

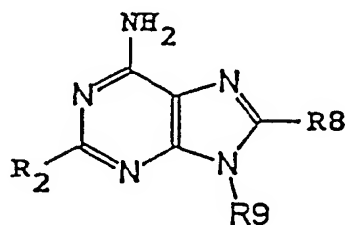


wherein R6 and alkyl have above mentioned meanings, and so obtained product is then optionally, either progressively or simultaneously, subjected to a nucleophilic substitution in positions 2 and 8 in order to replace the chlorine atom in position 2 and the bromine atom in position 8 by another substituents R2 and R8 that have above mentioned meanings.

The invention also relates to a method for preparing substituted nitrogen heterocyclic derivatives of formula I, wherein R6 is a halogen or hydrogen, R2, R8 and R9 have above mentioned meanings, A is a group of formula



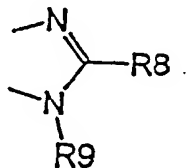
and Z is N, characterized in that a derivative of formula XVIII,



(XVIII)

wherein R2, R8 and R9 have above mentioned meanings, is halogenated *via* diazotation with using, for example, amyl nitrite/CH₂Br₂ or amyl nitrite/CHI₃, and so obtained derivative wherein R6 is halogen is optionally hydrogenolyzed with using H₂/Pd catalyst to obtain a corresponding derivative wherein R6 is hydrogen.

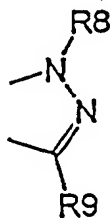
The invention also relates to a method for preparing substituted nitrogen heterocyclic derivatives of formula I, wherein R9 has above mentioned meanings and one substituent of R2, R6 or R8 is alkynyl or alkenyl containing 1 to 6 carbon atoms whereas two other substituents of R2, R6 and R8 have above mentioned meanings, A is a group of formula



and Z is N or CH, characterized in that a derivative of formula I, wherein R9 has above

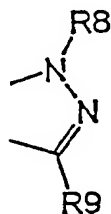
mentioned meanings and one substituent of R₂, R₆ and R₈ is a halogen whereas two other substituents of R₂, R₆ and R₈ have above mentioned meanings, is alkynylated in the position of the halogen with using alkyne/triphenylphosphine-PdCl₂/CuI/triethylamine system and so obtained 2-, 6- or 8-alkynyl derivative is optionally converted to 2-, 6- or 8-alkenyl derivative with using a Lindlar catalyst.

The invention also relates to a method for preparing substituted nitrogen heterocyclic derivatives of formula I, wherein R₂, R₆, R₈ and R₉ have above mentioned meanings, A is a group of formula



and Z is N, characterized in that 2,5-dialkyl-3-alkoxykarbonyl-4-aminopyrazole, wherein alkyls and alkoxy have above mentioned meanings, is reacted with formamidine acetate/triethylamine system to obtain 1,3-dialkyl-7-hydroxypyrazolo[4,3-d]pyrimidine and so obtained 7-hydroxy derivative is then optionally transferred to a corresponding 7-chloro derivative by reaction with thionylchloride, which product is optionally substituted to obtain a corresponding R₆-derivative, wherein R₆ has above mentioned meanings.

The invention also relates to a method for preparing substituted nitrogen heterocyclic derivatives of formula I, wherein R₂, R₆, R₈ and R₉ have above mentioned meanings, A is a group of formula



and Z is N, characterized in that 3,7-dialkylpyrazolo[4,3-d]pyrimidine, wherein alkyls have above mentioned meanings, is alkylated with using an appropriate alkylating agent in a system selected from a group consisting of K₂CO₃/dimethylformamide and Cs₂CO₃/dimethylformamide to obtain a mixture of trialkylated and tetraalkylated products and the trialkylated product is separated by means of the column chromatography.

The invention also relates to substituted nitrogen heterocyclic derivatives of formula I and pharmaceutically acceptable salts thereof for use as medicaments.

The invention also relates to substituted nitrogen heterocyclic derivatives of formula I for use as means for preparing affinity adsorption matrices, immobilized
5 enzymes for process control, immunoassay reagents, diagnostic samples, ^{14}C -, ^3H -,
avidin- or biotin-labelled compounds, oligonucleotides and diagnostic samples.

The invention also relates to substituted nitrogen heterocyclic derivatives of formula I for use as antimitotic drugs, in particular drugs for elimination or reduction of
10 viral spread or growth in tissue culture systems during the production of
biopharmaceutical or other products such as proteins and vaccines, drugs for elimination
or reduction of viral spread and growth in clinical samples such as blood, and drugs for
stopping of growth of tissue culture cells while leaving the cells to carry on with protein
production.

The invention also relates to a pharmaceutical composition with cytostatic,
15 anticancer, antimitotic, antineurodegenerative, immunosuppressive and antimicrobial
activity, characterised in that it comprises, in addition to auxiliary pharmaceutical
matters, at least one substituted nitrogen heterocyclic derivative of formula I.

The invention also relates to a combined pharmaceutical composition with
cytostatic effect, characterised in that it comprises, in addition to auxiliary
20 pharmaceutical matters, a combination of at least one substituted nitrogen heterocyclic
derivative of formula I and a cytostatic agent selected from a group consisting of
mitoxantron, cis- and carbo-platin, methotrexate, taxol and doxorubicin.

The invention also relates to a use of substituted nitrogen heterocyclic
derivatives of formula I for preparing medicaments destined for treating tumours,
25 cancers, psoriasis, rheumatoid arthritis, lupus, type I diabetes, multiple sclerosis,
restenosis, polycyclic kidney disease, host graft disease and gout, parasitoses, such as
those caused by fungi or protozoa, or Alzheimer's disease, or for preparing
antineurodegenerative drugs and suppress immunostimulation agents.

The novel compounds of this invention have a wide variety of diagnostic,
30 therapeutic and industrial utilities.

The compounds of this invention are suitable as intermediates for use in the
preparation of affinity absorption matrices. For example, the phosphonate groups in

matrix bound form are useful in the chromatographic separation of positively charged molecules. Other immobilised examples of the compounds herein are useful in purifying proteins, e.g., cell cycle enzymes (cdk's), enzymes involved in recognition of the compound of this invention, e.g. transport proteins. Suitable methods of incorporation of the compounds of this invention into polymeric resins will be readily apparent to the skilled artisan, for instance the compounds are incorporated by cross-linking hydroxyl groups of the phosphonate or hydroxymethyl substituents using cross-linking agents heretofore known. Linking through a group other than the heterocyclic base will produce a resin useful in hydrophobic affinity chromatography. Other suitable linking methods are described in Cihlar (supra).

The compounds of the formula I and their pharmaceutically acceptable salts inhibit selectively the enzyme p34^{cdc2}/cyclin B kinase and related cdks (cdk2, cdk5, cdk7, erk1, erk2).

In another embodiment, this invention is a method for inhibiting cdks and cell proliferation in mammals comprising administering a therapeutically effective amount of the composition of claim 1 to the mammal. The cdk inhibiting molecules are useful for treating cell proliferation disorders such as rheumatoid arthritis, lupus, type I diabetes, multiple sclerosis, cancer, restenosis, Alzheimer's disease, growth of parasites (animal, protists), host graft disease, and gout.

In another embodiment, this invention is a composition useful for treating fungal infections (fungi) in humans, animal, and in plants.

2,6,8,9-Tetrasubstituted and 6,8,9- or 2,8,9- trisubstituted adenine derivatives exhibit extremely high potency against DNA viruses on the part of the defined compounds. Such compounds otherwise have been considered to have little or no activity against DNA viruses. Moreover, surprisingly the chirally enriched or pure (S)-enantiomer is antivirally active. Heretofore, only the (R)-enantiomer was notably antivirally active, and then only against retroviruses.

In addition to other cdc2-related kinases, this kinase controls certain steps of cell division cycles, in particular the transition from G1 phase into the S phase and in particular the transition from the G2 phase into the M phase. Out the basis of this findings, it can be expected that the compounds of the formula I, II and their pharmaceutically acceptable salts can be used as antimitotic compounds or for treatment

of hypoproliferative diseases, such as cancer, restenosis and Alzheimer's disease. Thus in very low concentration (micromolar and lower), they are capable of inhibiting cell cycle transitions (G1/S, G2/M, M-phase/metaphase) carried out on the different animal bodies and embryos. Furthermore, the compounds are useful in treating autoimmune diseases, e.g. rheumatoid arthritis, lupus, type I diabetes, multiple sclerosis, etc., in treating of cancer, cardiovascular disease such as restenosis, host vs graft disease, gout, polycystic kidney disease and other proliferative diseases whose pathogenesis involves abnormal cell proliferation.

This invention also concerns novel compounds that have been discovered to be potent and specific inhibitors of I κ B- α kinase which prevents signal induced NF- κ B activation and cytokine synthesis *in vitro* and *in vivo*. Such inhibitors are expected to inhibit synthesis of cytokines and adhesion proteins whose synthesis is transcriptionally regulated by NF- κ B. Pro-inflammatory cytokines such as IL-1, IL-6, TNF and adhesion proteins (e.g. ICAM, VCAM and selections) belong to this class of molecules and have implicated in the pathogenesis of inflammatory diseases. Thus a potent inhibitor of I κ B- α kinase is useful in the clinical management of diseases where the NF- κ B activation is required for disease induction.

It also relates to novel compounds activating p53, the mammal cell's own natural brake gene for stopping uncontrolled cell proliferation (cancer), thus being able to switch off the cancer. p53 as well as retinoblastoma (Rb) are two well-characterised tumour suppressors whose inactivation may led to uncontrolled cell proliferation and malignancy. Phosphorylation of these two proteins, which are involved in the cell cycle regulatory mechanisms, is known to modulate their function. Thus a potent cdk inhibitor represent a good toll for treatment of cancers due to induction of wild type p53 protein in cancers expressing mutant p53.

In addition, studies carried out on the derivatives of the invention have demonstrated strong effect on apoptosis of many cancer cell lines. It has been seen that apoptosis can be induced at stage G1 or G2 and following damage of the DNA, some cells stop at stage G1 and p53-dependent apoptotic pathway is then induced. In other situations, it seems that cells stop at G2/M stage in response to damage caused to the DNA, and activation of an independent p53 apoptotic path is observed. This path has proved to be particularly significant in the therapy of tumours in which a less of active

p53 is observed. The interest is therefore assessed by application of the derivatives of the invention for stimulating on of p53-independent apoptosis in the cells, which have stopped at stage G2 through damage to the DNA using agents such as mitoxantrone or cis-platinum. The cdk inhibitors of this invention can thus increase the therapeutic potential of the anti-tumour agents currently used.

The compounds of this invention will generally be terminally incorporated into the oligonucleotide. If they do not contain phosphonyl group attached to the hydroxyl group, they optionally are incorporated internally into the sequence of the oligonucleotide. Terminally incorporated diphosphonyl compounds of this invention which contain no free hydroxyl capable of participating in chain elongation also are useful in DNA sequencing in essentially the same manner as deoxyNTPs have been used in the past (see example 8 of U.S. Patent 5,276,143). The nucleotide analogues of the invention (when diphosphorylated) are useful as chain terminators for dideoxynucleotide-type DNA sequencing protocols, provided that the nucleotide analogue lacks a free hydroxyl group suitable for polymerase mediated chain elongation. These compounds will not have R=hydroxymethyl and do not possess a cyclic structure incorporating the phosphorus atom (although compounds having such excluded structures can be intermediates). The nucleotide analogue is included in a kit with other reagents (such as Klenow polymerase or T4 polymerase, dNTPs, etc) needed for DNA sequencing (Otvos, et al. "Nucl. Acids.Res." 15:1763-1777 (1987)).

If the oligonucleotide-incorporated compound of this invention is binding-competent for its complementary sequence, i.e., if it is capable of base pairing, then this nucleotide monomer will participate in hybridisation. It is not necessary, however, that the incorporated nucleotide analogue of this invention base pair or otherwise participate in hybridisation. If it is located at the terminus of the oligonucleotide it will be useful as an immunological recognition site, or haptenic recognition site, to facilitate detection of the oligonucleotide by an antibody capable of binding the compound of this invention.

The compounds of this invention also are useful as linkers or spacers in preparation affinity absorption matrices (as opposed to functioning as affinity moieties per se as noted above), immobilised enzymes for process control, or immunoassay reagents. The compounds herein contain a multiplicity of functional groups that are suitable as sites for cross-linking desired substances. For example, it is conventional to

link affinity reagents such as hormones, peptides, antibodies, drugs, and the like to insoluble substrates. These insolubilised reagents are employed in known fashion to absorb binding partners for the affinity reagents from manufactured preparations, diagnostic samples and other impure mixture. Similarly, immobilised enzymes are used
5 to perform catalytic conversions with facile recovery of enzyme. Bifunctional compounds are commonly used to link analytes to detectable groups in preparing diagnostic reagents.

Many functional groups present in the compounds of this invention are suitable for use in cross-linking. For example, the phosphonic acid is used to form esters with
10 alcohols or amides with amines. The R groups substituted with OH, azido (which is reduced to amino if desired before cross-linking) or vinyl are exemplary suitable sites. Similarly, the amino, halo, acyl and other reactive sites found on group B are suitable. Suitable protection of reactive groups will be used where necessary while assembling the cross-linked reagent. In general, the compounds here are used by linking them through
15 phosphonic acid or amino group to the hydroxyl or amino groups of the linking partner in the same fashion as shown herein, and covalently bonded to the other binding partner through an R group. For example a first binding partner such as a steroid hormone is esterified and then this conjugate is cross-linked through hydroxymethyl R to cyanogen bromide activated Sepharose, whereby immobilised steroid is obtained. Other
20 chemistries for conjugation are well known. See for example Maggio, "Enzyme-Immunoassay" (CRC, 1988, pp 71-135) and references cited therein.

The oligonucleotides of this invention are labelled with any conventional detectable label, e.g. a fluorescent moiety such as fluorescein, radioisotopes such as ^{14}C or ^3H , stable free radicals, avidin, biotin and the like all of which previously have been used
25 as labels for immunoassays or diagnostic probes. The label will be present on the oligonucleotide or on the residue of an analogue of this invention. Suitable labelling methods are well known and are readily used with reactive groups such as hydroxyl, allyl and the like. A simple method is to label the compound of this invention with H_3 by proton exchange. The compounds also are biotinylated using conventional methods. See
30 for instance U.S. Patent 5,276,143 for analogous structures. However, the compounds of this invention also are useful directly in diagnostic probe assays without an exogenous detectable label. In one embodiment of this alternative, antibodies are raised against the

compounds of this invention. Such antibodies (which in turn are labelled or used in a double antibody configuration) bind to the analogue of this invention and thereby are useful in detecting its presence as label for a protein or oligonucleotide.

The compounds of the invention are useful for treatment of microbial infections, for treatment of tumours or for other indications described below. Microbial infections treatable by the compounds of this invention include viruses, parasites, yeast and fungi, but it is believed that the compounds are most effective against viruses, which constitutes the preferred utility. Exemplary viral infections include infections caused by DNA or RNA viruses including herpes viruses (CMV, HSV 1, HSV 2, EBV, varicella zoster virus (VZV), bovid herpesvirus type 1, equid herpesvirus type 1, HHV-6, papillomaviruses (HPV types 1-55 including carcinogenic HPV), flaviviruses (including yellow fever virus, African swine fever virus and Japanese encephalitis virus), togaviruses (including Venezuelan equine encephalomyelitis virus), influenza viruses (types A-C), retroviruses (HIV-1, HIV-2, HTLV-I, HTLV-II, SIV, FeLV, FIV, MoMSV), adenoviruses (types 1-8), poxviruses (vaccinia virus), enteroviruses (poliovirus types 1-3, Coxsackie, hepatitis A virus, and ECHO virus), gastroenteritis viruses (Norwalk viruses, rotaviruses), hantaviruses (Hantaan virus), polyomavirus, papovaviruses, rhinoviruses, parainfluenza virus types 1-4, rabies virus, respiratory syncytial virus (RSV), hepatitis viruses A, B, C and E, and the like.

The antiviral activity of individual compounds is determined by routine assay of antiviral (or other antimicrobial) activity using enzyme inhibition assays, tissue culture assays, animal model assays and the like as will be understood by those skilled in the art.

Protozoan parasite infections are treated using the compounds of the invention. The term protozoa include those members of the subphyla *Sarcomastigophora* and *Sporozoa* of the phylum *Protozoa*. More particularly, the term protozoa as used herein include genera of parasitic protozoa, which are important to man, because they either cause disease in man or in his domestic animals. These genera for the most part are classified in the superclass *Mastigophora* of the subphylum *Sarcomastigophora* and the class *Telesporea* of the subphylum *Sporozoa* in the classification according to Baker (1969). Illustrative genera of these parasitic protozoa include *Histomonas*, *Pneumocystis*, *Trypanosoma*, *Giardia*, *Trichomonas*, *Eimeria*, *Isopora*, *Leishmania*, *Entamoeba*, *Toxoplasma* and *Plasmodium*. Parasitic protozoans include *Plasmodium falciparum*,

Plasmodium berghei, *Plasmodium malariae*, *Plasmodium vivax*, *Leishmania braziliensis*, *Leishmania donovani*, *Trypanosoma cruzi*, *Trypanosoma brucei*, *Trypanosoma rhodesiense*, *Pneumocystis carinii*, *Entamoeba histolytica*, *Trichomonas vaginalis* and the like (de Vries, E., et al, "Mol. Biochem. Parasitol" 47:43-50 (1991) and trypanosomes (Kaminsky et al. "J.Parasitol." 80(6): 1026-30 (1994). The compounds in which R is CH₂OH and B is 3-deazaadenine are particularly interesting in the treatment of malarial parasites.

Compounds of the invention are used to treat yeast or fungal infections caused by *Candida glabrata*, *Candida tropicalis*, *Candida albicans*, and other *Candida* species, *Cryptococcus* species including *Cryptococcus neoformans*, *Blastomyces* species including *Blastomyces dermatidis*, *Torulopsis* species including *Torulopsis glabrata*, *Coccidioides* species including *Coccidioides immitis*, *Aspergillus* species and the like.

The compounds of the invention can also be (1) applied to tissue culture systems to eliminate or reduce viral spread or growth during the production of biopharmaceutical or other products (such as proteins or vaccines), (2) used to eliminate or reduce viral spread or growth in clinical sample (such as blood), and (3) used to stop growth of tissue culture cells while leaving the cells to carry on with protein production.

The compounds herein have been found to suppress immunostimulation. Accordingly, they can suppress metabolic activities of T-lymphocytes stimulated by diverse agents, e.g. concavalin A, they principally will find application in the treatment of autoimmune diseases, e.g. arthritis, or in suppression of transplant rejection. Their therapeutically active concentrations are in the range of 1 mg/kg to 50 mg/kg of body weight.

25 Therapeutic administration

Suitable routes for administration include oral, rectal, vassal, topical (including ocular, buccal and sublingual), vaginal and parental (including subcutaneous, intramuscular, intravitreal, intravenous, intradermal, intrathecal and epidural). The preferred route of administration will depend upon the condition of the patient, the toxicity of the compound and the site of infection, among other considerations known to the clinician.

The therapeutical composition comprise about 1% to about 95% of the active ingredient, single-dose forms of administration preferably comprising about 20% to about 90% of the active ingredient and administration forms which are not single-dose preferably comprising about 5% to about 20% of the active ingredient. Unit dose forms are, for example, coated tablets, tablets, ampoules, vials, suppositories or capsules. Other forms of administration are, for example, ointments, creams, pastes, foams, tinctures, lipsticks, drops, sprays, dispersions and the like. Examples are capsules containing from about 0.05 g to about 1.0 g of the active ingredient.

The pharmaceutical compositions of the present invention are prepared in a manner known per se, for example by means of conventional mixing, granulating, coating, dissolving or lyophilising processes.

Preferably, solutions of the active ingredient, and in addition also suspensions or dispersions, especially isotonic aqueous solutions, dispersions or suspensions, are used, it being possible for these to be prepared before use, for example in the case of lyophilised compositions which comprise the active substance by itself or together with a carrier, for example mannitol. The pharmaceutical compositions can be sterilised and/or comprise excipients, for example preservatives, stabilisers, wetting agents and/or emulsifiers, solubilizing agents, salts for regulating the osmotic pressure and/or buffers, and they are prepared in a manner known per se, for example by means of conventional dissolving or lyophilising processes. The solutions or suspensions mentioned can comprise viscosity-increasing substances, such as sodium carboxymethylcellulose, carboxymethylcellulose, dextran, polyvinylpyrrolidone or gelatin.

Suspensions in oil comprise, as the oily component, the vegetable, synthetic or semisynthetic oils customary for injection purposes. Oils which may be mentioned are, in particular, liquid fatty acid esters which contain, as the acid component, a long-chain fatty acid having 8 - 22, in particular 12-22, carbon atoms, for example lauric acid, tridecylic acid, myristic acid, pentadecylic acid, palmitic acid, margaric acid, stearic acid, arachidinic acid, behenic acid or corresponding unsaturated acids, for example oleic acid, elaidic acid, euric acid, brasidic acid or linoleic acid, if appropriate with the addition of antioxidants, for example vitamin E, β -carotene or 3,5-di-tert-butyl-4-hydroxytoluene. The alcohol component of these fatty acid esters has not more than 6 carbon atoms and is mono- or polyhydric, for example mono-, di- or trihydric alcohol,

for example methanol, ethanol, propanol, butanol, or pentanol, or isomers thereof, but in particular glycol and glycerol. Fatty acid esters are therefore, for example: ethyl oleate, isopropyl myristate, isopropyl palmitate, "Labrafil M 2375" (polyoxyethylene glycerol trioleate from Gattefoscé, Paris), "Labrafil M 1944 CS" (unsaturated polyglycolated glycerides prepared by an alcoholysis of apricot kernel oil and made up of glycerides and polyethylene glycol esters; from Gattefoscé, Paris), "Labrasol" (saturated polyglycolated glycerides prepared by an alcoholysis of TCM and made up of glycerides and polyethylene glycol esters; from Gattefoscé, Paris) and/or "Miglyol 812" (triglyceride of saturated fatty acids of chain length C_8 to C_{12} from Hüls AG, Germany), and in particular vegetable oils, such as cottonseed oil, almond oil, olive oil, castor oil, sesame oil, soybean oil and, in particular, groundnut oil.

The preparation of the injection compositions is carried out in the customary manner under sterile conditions, as are bottling, for example in ampoules or vials, and closing of the containers.

For example, pharmaceutical compositions for oral use can be obtained by combining the active ingredient with one or more solid carriers, if appropriate granulating the resulting mixture, and, if desired, processing the mixture or granules to tablets or coated tablet cores, if appropriate by addition of additional excipients.

Suitable carriers are, in particular, fillers, such as sugars, for example lactose, sucrose, mannitol or sorbitol, cellulose preparations and/or calcium phosphates, for example tricalcium phosphate, or calcium hydrogen phosphate, and furthermore binders, such as starches, for example maize, wheat, rice or potato starch, methylcellulose, hydroxypropylmethylcellulose, sodium carboxymethylcellulose and/or polyvinylpyrrolidone, and/or, if desired, disintegrators, such as the above mentioned starches, and furthermore carboxymethyl-starch, crosslinked polyvinylpyrrolidone, alginic acid or a salt thereof, such as sodium alginate. Additional excipients are, in particular, flow regulators and lubricants, for example salicylic acid, talc, stearic acid or salts thereof, such as magnesium stearate or calcium stearate, and/or polyethylene glycol, or derivatives thereof.

Coated tablet cores can be provided with suitable coatings which, if appropriate, are resistant to gastric juice, the coatings used being, inter alia, concentrated sugar solutions, which, if appropriate, comprise gum arabic, talc, polyvinylpyrrolidone,

polyethylene glycol and/or titanium dioxide, coating solutions in suitable organic solvents or solvent mixtures or, for the preparation of coatings which are resistant to gastric juice, solutions of suitable cellulose preparations, such as acetylcellulose phthalate or hydroxypropylmethylcellulose phthalate. Dyes or pigments can be admixed
5 to the tablets or coated tablet coatings, for example for identification or characterisation of different doses of active ingredient.

Pharmaceutical compositions, which can be used orally, are also hard capsules of gelatin and soft, closed capsules of gelatin and a plasticiser, such as glycerol or sorbitol. The hard capsules can contain the active ingredient in the form of granules, mixed for
10 example with fillers, such as maize starch, binders and/or lubricants, such as talc or magnesium stearate, and stabilisers if appropriate. In soft capsules, the active ingredient is preferably dissolved or suspended in suitable liquid excipients, such as greasy oils, paraffin oil or liquid polyethylene glycols or fatty acid esters of ethylene glycol or propylene glycol, it being likewise possible to add stabilisers and detergents, for
15 example of the polyethylene sorbitan fatty acid ester type.

Other oral forms of administration are, for example, syrups prepared in the customary manner, which comprise the active ingredient, for example, in suspended form and in a concentration of about 5% to 20%, preferably about 10% or in a similar concentration which results in a suitable individual dose, for example, when 5 or 10 ml
20 are measured out. Other forms are, for example, also pulverulent or liquid concentrates for preparing of shakes, for example in milk. Such concentrates can also be packed in unit dose quantities.

Pharmaceutical compositions, which can be used rectally, are, for example, suppositories that comprise a combination of the active ingredient with a suppository
25 base. Suitable suppository bases are, for example, naturally occurring or synthetic triglycerides, paraffin hydrocarbons, polyethylene glycols or higher alkanols.

Compositions which are suitable for parenteral administration are aqueous solutions of an active ingredient in water-soluble form, for example of water-soluble salt, or aqueous injection suspensions, which comprise viscosity-increasing substances, for
30 example sodium carboxymethylcellulose, sorbitol and/or dextran, and if appropriate stabilisers. The active ingredient can also be present here in the form of a lyophilisate, if appropriate together with excipients, and be dissolved before parenteral administration

by addition of suitable solvents. Solutions such as are used, for example, for parental administration can also be used as infusion solutions. Preferred preservatives are, for example. Antioxidants, such as ascorbic acid, or microbicides, such as sorbic or benzoic acid.

5 Ointments are oil-in-water emulsions, which comprise not more than 70%, but preferably 20 - 50% of water or aqueous phase. The fatty phase consists, in particular, hydrocarbons, for example vaseline, paraffin oil or hard paraffin's, which preferably comprise suitable hydroxy compounds, such as fatty alcohol's or esters thereof, for example cetyl alcohol or wool wax alcohols, such as wool wax, to improve the water-
10 binding capacity. Emulsifiers are corresponding lipophilic substances, such as sorbitan fatty acid esters (Spans), for example sorbitan oleate and/or sorbitan isostearate. Additives to the aqueous phase are, for example, humectants, such as polyalcohols, for example glycerol, propylene glycol, sorbitol and/or polyethylene glycol, or preservatives and odoriferous substances.

15 Fatty ointments are anhydrous and comprise, as the base, in particular, hydrocarbons, for example paraffin, vaseline or paraffin oil, and furthermore naturally occurring or semisynthetic fats, for example hydrogenated coconut-fatty acid triglycerides, or, preferably, hydrogenated oils, for example hydrogenated groundnut or castor oil, and furthermore fatty acid partial esters of glycerol, for example glycerol
20 mono- and/or distearate, and for example, the fatty alcohols. They also contain emulsifiers and/or additives mentioned in connection with the ointments which increase uptake of water.

Creams are oil-in-water emulsions, which comprise more than 50% of water. Oily bases used are, in particular, fatty alcohols, for example lauryl, cetyl or stearyl
25 alcohols, fatty acids, for example palmitic or stearic acid, liquid to solid waxes, for example isopropyl myristate, wool wax or beeswax, and/or hydrocarbons, for example vaseline (petrolatum) or paraffin oil. Emulsifiers are surface-active substances with predominantly hydrophilic properties, such as corresponding nonionic emulsifiers, for example fatty acid esters of polyalcohols or ethyleneoxy adducts thereof, such as
30 polyglyceric acid fatty acid esters or polyethylene sorbitan fatty esters (Tweens), and furthermore polyoxyethylene fatty alcohol ethers or polyoxyethylene fatty acid esters, or corresponding ionic emulsifiers, such as alkali metal salts of fatty alcohol sulfates, for

example sodium lauryl sulfate, sodium cetyl sulfate or sodium stearyl sulfate, which are usually used in the presence of fatty alcohols, for example cetyl stearyl alcohol or stearyl alcohol. Additives to the aqueous phase are, inter alia, agents which prevent the creams from drying out, for example polyalcohols, such as glycerol, sorbitol, propylene glycol
5 and/or polyethylene glycols, and furthermore preservatives and odoriferous substances.

Pastes are creams and ointments having secretion-absorbing powder constituents, such as metal oxides, for example titanium oxide or zinc oxide, and furthermore talc and/or aluminium silicates, which have the task of binding the moisture or secretions present.

10 Foams are administered from pressurised containers and they are liquid oil-in-water emulsions present in aerosol form. As the propellant gases, halogenated hydrocarbons, such as chlorofluoro-lower alkanes, for example dichlorofluoromethane and dichlorotetrafluoroethane, or, preferably, non-halogenated gaseous hydrocarbons, air, N₂O, or carbon dioxide are used. The oily phases used are, inter alia, those
15 mentioned above for ointments and creams, and the additives mentioned there are likewise used.

Tinctures and solutions usually comprise an aqueous-ethanolic base to which, humectants for reducing evaporation, such as polyalcohols, for example glycerol, glycols and/or polyethylene glycol, and re-oiling substances, such as fatty acid esters
20 with lower polyethylene glycols, i.e. lipophilic substances soluble in the aqueous mixture to substitute the fatty substances removed from the skin with the ethanol, and, if necessary, other excipients and additives, are admixed.

The present invention further provides veterinary compositions comprising at least one active ingredient as above defined together with a veterinary carrier therefor.
25 Veterinary carriers are materials for administering the composition and may be solid, liquid or gaseous materials, which are inert or acceptable in the veterinary art and are compatible with the active ingredient. These veterinary compositions may be administered orally, parenterally or by any other desired route.

The invention also relates to a process or method for treatment of the disease
30 states mentioned above. The compounds can be administered prophylactically or therapeutically as such or in the form of pharmaceutical compositions, preferably in an amount, which is effective against the diseases mentioned. With a warm-blooded animal,

for example a human, requiring such treatment, the compounds are used, in particular, in the form of pharmaceutical composition. A daily dose of about 0.1 to about 5 g, preferably 0.5 g to about 2 g, of a compound of the present invention is administered here for a body weight of about 70 kg.

5

Examples of the invention

EXAMPLE 1:

2-(3-hydroxypropylamino)-6-benzylamino-8-bromo-9-isopropylpurine.

- 10 1 mmol of 2-(3-hydroxypropylamino)-6-benzylamino-9-isopropylpurine was dissolved in chloroform and 1.1 mmol of bromine was added. The 8-bromo derivative hydrobromide was removed by filtration and crystallized from n-propanol - ether. M.p. 156-161°C. Yield 90 %. TLC: chloroform-methanol (96 : 4), single spot. MS-ESI(+) m/z: 420, 422 [M+H]⁺

15

Table 1 Compounds Prepared by the Method of Example 1

SUBSTITUENT			
C2	N6	C8	N9
2-hydroxyethylamino	benzylamino	bromo	methyl
3-hydroxypropylamino	benzylamino	bromo	isopropyl
(R)-1-(hydroxymethyl) propylamino	benzylamino	bromo	isopropyl
(R)-1-(hydroxymethyl) propylamino	3-iodobenzylamino	bromo	isopropyl
diethanolamino	3-chloroanilino	bromo	isopropyl
(R)-1-(hydroxymethyl)isobutylamino	4-methoxybenzylamino	bromo	isopropyl
(R)-1-(hydroxymethyl)isobutylamino	3-chloroanilino	bromo	isopropyl

EXAMPLE 2:

2-(3-hydroxypropylamino)-6-benzylamino-8-methylthio-9-isopropylpurine.

- 20 2-(3-Hydroxypropylamino)-6-benzylamino-8-bromo-9-isopropylpurine (0.15 mmol), prepared as described in **Example 1**, was dissolved in 2 mL dimethylformamide, 0.9 mmol of CH₃SNa was added and the mixture stirred at 40°C for 1 hour. The solvent was removed *in vacuo* and the rest partitioned between water-EtOAc. The organic layer was

dried, evaporated and the rest purified by column chromatography (silica gel, chloroform-heptane (8 : 2)). Yield 75 %. M. p. 64-67 °C.

¹H NMR (300 MHz, CDCl₃): 7.28-740 m, 5.55 t, 4.85 t, 4.75 d, 4.65 m, 3.62 m, 2.62 s, 1.74 dd, 1.60 d.

5 MS-ESI(+) m/z: 387 [M+H]⁺

Table 2 Compounds Prepared by the Method of Example 2

SUBSTITUENT			
C2	N6	C8	N9
2-hydroxyethylamino	benzylamino	methyl	methyl
2-hydroxyethylamino	benzylamino	mercapto	methyl
2-hydroxyethylamino	benzylamino	hydroxy	methyl
2-hydroxyethylamino	benzylamino	amino	methyl
2-hydroxyethylamino	benzylamino	2-hydroxyethylamino	methyl
2-hydroxyethylamino	benzylamino	aminomethylamino	methyl
3-hydroxypropylamino	benzylamino	methyl	isopropyl
3-hydroxypropylamino	benzylamino	mercapto	isopropyl
3-hydroxypropylamino	benzylamino	hydroxy	isopropyl
3-hydroxypropylamino	benzylamino	amino	isopropyl
3-hydroxypropylamino	benzylamino	2-hydroxyethylamino	isopropyl
3-hydroxypropylamino	benzylamino	aminomethylamino	isopropyl
(R)-1-(hydroxymethyl) propylamino	benzylamino	methyl	isopropyl
(R)-1-(hydroxymethyl) propylamino	benzylamino	mercapto	isopropyl
(R)-1-(hydroxymethyl) propylamino	benzylamino	hydroxy	isopropyl
(R)-1-(hydroxymethyl) propylamino	benzylamino	amino	isopropyl
(R)-1-(hydroxymethyl) propylamino	benzylamino	2-hydroxyethylamino	isopropyl
(R)-1-(hydroxymethyl) propylamino	benzylamino	aminomethylamino	isopropyl
(R)-1-(hydroxymethyl) propylamino	3-iodobenzylamino	methyl	isopropyl
(R)-1-(hydroxymethyl) propylamino	3-iodobenzylamino	mercapto	isopropyl
(R)-1-(hydroxymethyl) propylamino	3-iodobenzylamino	hydroxy	isopropyl
(R)-1-(hydroxymethyl) propylamino	3-iodobenzylamino	amino	isopropyl
(R)-1-(hydroxymethyl) propylamino	3-iodobenzylamino	2-hydroxyethylamino	isopropyl
(R)-1-(hydroxymethyl) propylamino	3-iodobenzylamino	aminomethylamino	isopropyl
diethanolamino	3-chloroanilino	methyl	isopropyl
diethanolamino	3-chloroanilino	mercapto	isopropyl

diethanolamino	3-chloroanilino	hydroxy	isopropyl
diethanolamino	3-chloroanilino	amino	isopropyl
diethanolamino	3-chloroanilino	2-hydroxyethylamino	isopropyl
diethanolamino	3-chloroanilino	aminomethylamino	isopropyl
(R)-1-(hydroxymethyl)isobutylamino	4-methoxybenzylamino	mercapto	isopropyl
(R)-1-(hydroxymethyl)isobutylamino	4-methoxybenzylamino	hydroxy	isopropyl
(R)-1-(hydroxymethyl)isobutylamino	4-methoxybenzylamino	amino	isopropyl
(R)-1-(hydroxymethyl)isobutylamino	4-methoxybenzylamino	2-hydroxyethylamino	isopropyl
(R)-1-(hydroxymethyl)isobutylamino	4-methoxybenzylamino	aminomethylamino	isopropyl
(R)-1-(hydroxymethyl)isobutylamino	3-chloroanilino	methyl	isopropyl
(R)-1-(hydroxymethyl)isobutylamino	3-chloroanilino	mercapto	isopropyl
(R)-1-(hydroxymethyl)isobutylamino	3-chloroanilino	hydroxy	isopropyl
(R)-1-(hydroxymethyl)isobutylamino	3-chloroanilino	amino	isopropyl
(R)-1-(hydroxymethyl)isobutylamino	3-chloroanilino	2-hydroxyethylamino	isopropyl
(R)-1-(hydroxymethyl)isobutylamino	3-chloroanilino	aminomethylamino	isopropyl

EXAMPLE 3:

6-(3-Methoxybenzylamino)-2-chloro-8-bromo-9-isopropylpurine.

1 mmol of 6-(3-methoxybenzylamino)-2-chloro-9-isopropylpurine was brominated with
 5 1.1 mmol of bromine as described in **Example 1**. The crude 2,8-dihalogeno derivative prepared was alkalinized with NH_3/MeOH and extracted between water - EtOAc. The product crystallizes after partial evaporation of ethylacetate. M.p.123-125°C, yield 88 %. MS-ESI (+) m/z 409.5, 411.4, 413.3 $[\text{M}+\text{H}]^+$

EXAMPLE 4:

6-Phenylamino-2-(3-aminopropylamino)-9-isopropyl-8-aminopurine.

6-Phenylamino-2-(3-aminopropylamino)-9-isopropyl-9-benzylaminopurine, prepared from appropriate 8-bromo derivative, was dissolved in glacial acetic acid. To this solution, 50 mg Pd/BaSO_4 (10 %) was added and the mixture hydrogenated to constant
 15 consumption of hydrogen (2 hours). The catalyst was removed by centrifugation and acetic acid was evaporated *in vacuo*. The rest was purified by column chromatography (silica gel, chloroform-MeOH-conc. NH_4OH (95 : 5 : 1). Yield 75 %.

MS-ESI (+) m/z 340.5 $[\text{M}+\text{H}]^+$

EXAMPLE 5:

6-(4-Methoxybenzylamino)-2,8-bis(3-hydroxypropylamino)-9-isopropylpurine.

6-(4-Methoxybenzylamino)-2,8-bis(3-hydroxypropylamino)purine (0.07 mmol) was stirred in 2 mL dimethylformamide with 0.14 mmol of allylbromide and with 0.36 mmol of dry potassium carbonate at ambient temperature (6 hours). The solvent was evaporated to dryness and the rest extracted between ethylacetate - water. The organic layer was dried, ethylacetate was evaporated and the rest purified by column chromatography (silica gel, chloroform). Yield 55%. MS-ESI(+) m/z 441.4 [M+H]⁺

Table 3 Compounds Prepared by the Method of Example 5

SUBSTITUENT			
C2	N6	C8	N9
2-hydroxyethylamino	benzylamino	2-hydroxyethylamino	allyl
3-hydroxypropylamino	4-methoxybenzylamino	3-hydroxypropylamino	allyl
2-aminoethylamino	benzylamino	2-aminoethylamino	isopropyl
3-aminopropylamino	benzylamino	3-aminopropylamino	isopropyl
aminomethylamino	benzylamino	aminomethylamino	isopropyl
diethanolamino	benzylamino	diethanolamino	isopropyl
hydroxymethylamino	benzylamino	hydroxymethylamino	isopropyl
3-aminopropylamino	benzylamino	3-aminopropylamino	isopropyl
2-hydroxyethylamino	3-chloroanilino	2-hydroxyethylamino	isopropyl
3-hydroxypropylamino	3-chloroanilino	3-hydroxypropylamino	isopropyl
2-aminoethylamino	3-chloroanilino	2-aminoethylamino	isopropyl
3-aminopropylamino	3-chloroanilino	3-aminopropylamino	isopropyl
diethanolamino	benzylamino	diethanolamino	isopropyl
hydroxymethylamino	benzylamino	hydroxymethylamino	isopropyl
(R)-1-(hydroxymethyl)isobutylamino	3-chloroanilino	(R)-1-(hydroxymethyl)isobutylamino	isopropyl

EXAMPLE 6:

6-Benzylamino-8-hydroxyethylamino-9-isopropylpurine.

1 mmol of 6-chloropurine was alkylated with isopropylbromide in DMF as described in **Example 5**. The product, 6-chloro-9-isopropylpurine was brominated in acetic acid

similarly as described in **Example 1**. After purification by column chromatography, 6-benzylamino-9-isopropyl-8-bromopurine was treated with 2-aminoethanol to give 6-benzylamino-8-hydroxyethylamino-9-isopropylpurine. Yield 40 %.

MS-ESI(+) m/z 326 [M+H]⁺

5

EXAMPLE 7:

2-(3-Hydroxypropylamino)-9-methyl-8-(3-hydroxypropyloxy)purine.

6-Amino-2-(3-hydroxypropylamino)-9-methyl-8-(3-hydroxypropyloxy)purine was brominated in the position 6 with amylnitrite/CH₂Br₂. The 6-bromo derivative was then hydrogenolyzed with PdO/BaSO₄ in strong alkaline solution to give 2,8,9-trisubstituted purine. The crude product was purified by column chromatography (silica gel, chloroform-MeOH-conc. NH₄OH (8 : 2 : 0.2)). The total yield was 35 %. MS-ESI(+) 281 [M+H]⁺

15 **EXAMPLE 8:**

7-Hydroxy-1-methyl-3-isopropylpyrazolo[4,3]pyrimidine.

0.02 mmol of 2-Methyl-4-amino-3-alkoxycarbonyl-5-isopropylpyrazole (prepared from appropriate 4-nitro derivative by catalytic hydrogenation) was dissolved in 30 mL of 2-ethoxyethanol and 0.02 mmol of formamidin acetate and 5.2 mL of triethylamine were added. The mixture was heated to 90 °C for 2 hours. The solution was concentrated *in vacuo*, dissolved in chloroform to give crystalline product. After recrystallization from ethanol m.p. 295-298 °C.

MS-ESI(+) m/z 193 [M+H]⁺

25 **EXAMPLE 9:**

7-(2-Hydroxybenzylamino)-1-methyl-3-isopropylpyrazolo[4,3]pyrimidine.

1.1 mmol of 7-Hydroxy-1-methyl-3-isopropylpyrazolo[4,3]pyrimidine (**Example 8**) was dissolved in the mixture of 0.12 mL of dimethylformamide and 5 mL dry chloroform. 11 mmol of thionylchloride (0.81 mL) was added and the solution heated to 80 °C for 1 hour. A new portion of 0.8 mL of thionylchloride and more chloroform-dimethylformamide were then added and heating continued for 3 hours. The solvents were evaporated *in vacuo* and the rest partitioned between water-chloroform. The

30

organic extract was dried and used in next reaction without purification. The chloroform solution was heated with excess of 2-hydroxybenzylamine and N-ethyl-diisopropylamine for 1 hour. The product was purified by column chromatography (silica gel, chloroform-MeOH-AcOH (20 : 0.4 : 0.1)). M.p. 205-210 °C. Yield 40 %. MS-ESI(+) m/z 297

5 [M+H]⁺

EXAMPLE 10.

6-(Benzylamino)-2-[(1-hydroxymethyl)propylamino]-8-chloro-9-isopropylpurine.

6-(Benzylamino)-2,8-dichloropurine, prepared from 2,6,8-trichloropurine, m.p. 220-
10 223°C, 50 % yield, was dissolved in 2-aminobutanol and heated to 150°C for 12 hours. After evaporation of the solvent *in vacuo* the product, 6-benzylamino-2-[(1-hydroxymethyl)propylamino]-8-chloropurine was purified by column chromatography (silica gel, chloroform - MeOH - conc. NH₄OH 9 : 1 : 0.1) with total yield 36 %. MS-ESI(+) m/z 348 [M+H]⁺. The product was subsequently alkylated with
15 isopropyl iodide/K₂CO₃/dimethylformamide to give, after preparative column chromatography (silica gel, chloroform) 70 % yield of the desired compound. MS (ESI+): 388.4, 390.6 (M+H)⁺.

EXAMPLE 11: Preparation of Affinity Sorbent

20 Preparation of 2-(2-aminopropylamino)-6-(3-hydroxypropylamino)-8-bromo-9-isopropylpurine Epoxy activated Sepharose 6B Affinity Matrix.

Freeze-dried epoxy activated Sepharose 6B (Pharmacia LKB, Piscataway, NJ) was chosen for the coupling reaction due to its ability to form an ether bond between a hydroxyl-containing ligand and the epoxide group on the Sepharose. The gel was
25 swollen according to the manufacturer's instructions, (100 mg) of any one of the compound defined in claim 1 and 56 (preferably with R6 = aminooctylamino, 3- or 4-benzylamino, etc.) was dissolved in 1 ml coupling solution (1.2:1, v/v, DMF, 0.1N NaOH) and mixed with 0.5 ml of swollen gel at pH 10-11 for 72 h at room temperature with gentle agitation. Excess reactive groups were blocked with 1M ethanolamine for 4
30 hours at 50°C and the gel slurry was poured into 1-ml syringe column. The resin was activated with three alternating cycles of twenty column volumes each of pH 4.0 (0.1M acetate, 0.5 M NaCl) and pH 8.0 (0.1M tris-HCl, 0.5 M NaCl) buffers followed by

twenty column volumes of reaction buffer (20 mM HEPES, pH 7.3, 10 mM MgCl_2 , 15 mM glycerophosphate, 0.5 mM sodium orthovanadate, 0.5 mM EGTA). The column was stored at 4°C in the reaction buffer containing 0.1% sodium azide and regenerated prior to each use with alternating cycles of low and high pH as described above.

5 The Sf9 insect cell lysate (500 μg protein in 1-ml reaction buffer) was passed over the affinity column matrix sequentially five times and the flow through was saved (unbound material). The matrix was then washed three times with 1 ml reaction buffer (wash 1-3) then three times each reaction buffer containing 0.5M NaCl (eluate 1-3). The coupled proteins were eluted at low pH (pH 4.0, 0.1M acetate, 0.5M NaCl) as described
10 above and aliquots (20 μl from 1 ml) of each sample were assayed for their ability to phosphorylate histone H1 and other substrate proteins as described in Example 12. The presence of CDK complexes was also determined by SDS-PAGE.

EXAMPLE 12: CDK Inhibition Assays

15 Proteins

Cyclin-dependent kinases ($\text{p34}^{\text{cdc}2}$, $\text{p33}^{\text{cdk}2}$, $\text{p33}^{\text{cdk}4}$) and cyclins (cyclin B, E and D1) are produced in Sf9 insect cells coinfecting with appropriate baculoviral constructs. The cells are harvested 68-72 hrs post infection in lysis buffer for 30 min on ice and the soluble fraction is recovered by centrifugation at 14,000 g for 10 min. The protein
20 extract is stored at -80 °C.

Rb-GST is produced using an *E. coli* expression system, containing sequence encoding C terminus of retinoblastoma protein (aminoacids 773-928), which is known to be phosphorylated by $\text{p33}^{\text{cdk}4}$ kinase. The fusion protein is purified on glutathione-agarose beads.

25 Lysis buffer: 50mM Tris pH 7.4, 150mM NaCl, 5mM EDTA, 20mM NaF, 1%Tween, 1mM DTT, 0.1mM PMSF, leupeptine, aprotinin

Enzyme inhibition assays

To carry out experiments on kinetics under linear conditions, the final point test system for kinase activity measurement is used. The kinase is added to reaction mixture
30 in such a way as to obtain linear activity with respect to the concentration of enzyme and with respect to time.

The p34^{cdc2} and p33^{cdk2} kinase inhibition determination involves the use of 1mg/ml histone H1 (Sigma, type III-S) in the presence of 15μM [γ -³²P]ATP (500-100 cpm/pmol) (Amersham) in a final volume of 20 μl, inhibition of p33^{cdk4} kinase is determined with Rb-GST (0,2mg/ml) as a substrate. Kinase activity is determined at 30 °C in the kinase buffer.

Tested compounds are usually dissolved to 100mM solutions in DMSO, final concentration of DMSO in reaction mixture never exceeds 1%. The controls contain suitable dilutions of DMSO.

After 10 min, addition 3x SDS sample buffer stops the incubations. Phosphorylated proteins are separated electrophoretically using 12,5% SDS polyacrylamide gel. The measurement of kinase activity is done using digital image analysis.

The kinase activity is expressed as a percentage of maximum activity, the apparent inhibition constants are determined by graphic analysis. Kinase buffer: 50mM Hepes pH 7,4, 10mM MgCl₂, 5mM EGTA, 10mM 2-glycerolphosphate, 1mM NaF, 1mM DTT

Table 4 Kinase Inhibitory Activity of 2,6,8,9-Tetrasubstituted Purine Derivatives

SUBSTITUENT				CDC2	IκB-α
C2	N6	C8	N9	IC ₅₀ (μM)	IC ₅₀ (μM)
2-hydroxyethylamino OLOMOUCINE	benzylamino	-	methyl	7	15.4
2-hydroxyethylamino	benzylamino	fluoro	methyl	4.2	7.8
2-hydroxyethylamino	benzylamino	bromo	methyl	16.2	22.8
2-hydroxyethylamino	benzylamino	mercapto	methyl	22.4	35.7
2-hydroxyethylamino	benzylamino	hydroxy	methyl	13.5	16.2
2-hydroxyethylamino	benzylamino	amino	methyl	14.2	17.1
2-hydroxyethylamino	benzylamino	2-hydroxy-ethylamino	methyl	125.2	153.8
2-hydroxyethylamino	benzylamino	aminomethyl-amino	methyl	114.3	126.5
3-hydroxypropylamino BOHEMINE	benzylamino	-	isopropyl	1	3.2

3-hydroxypropylamino	benzylamino	bromo	isopropyl	12.93	24.5
3-hydroxypropylamino	benzylamino	mercapto	isopropyl	13.1	20.9
3-hydroxypropylamino	benzylamino	hydroxy	isopropyl	6.48	12.6
3-hydroxypropylamino	benzylamino	amino	isopropyl	8.52	12.8
3-hydroxypropylamino	benzylamino	2-hydroxy-ethylamino	isopropyl	143.8	152.6
3-hydroxypropylamino	benzylamino	aminomethyl-amino	isopropyl	123.5	138.7
(R)-1-(hydroxymethyl)-propylamino ROSCOVITINE	benzylamino		isopropyl	0.45	1.4
(R)-1-(hydroxymethyl)-propylamino	benzylamino	fluoro	isopropyl	0.38	1.2
(R)-1-(hydroxymethyl)-propylamino	benzylamino	bromo	isopropyl	10.38	11.2
(R)-1-(hydroxymethyl)-propylamino	benzylamino	mercapto	isopropyl	41.11	43.8
(R)-1-(hydroxymethyl)-propylamino	benzylamino	hydroxy	isopropyl	10.15	10.94
(R)-1-(hydroxymethyl)-propylamino	benzylamino	amino	isopropyl	20.17	20.85
(R)-1-(hydroxymethyl)-propylamino	benzylamino	2-hydroxy-ethylamino	isopropyl	151.93	164.95
(R)-1-(hydroxymethyl)-propylamino	benzylamino	aminomethyl-amino	isopropyl	181.05	193.28
(R)-1-(hydroxymethyl)-isobutylamino PURVALANOL A	3-chloroanilino		isopropyl	4 nM	11.2 nM

(R)-1-(hydroxymethyl)-isobutylamino	3-chloroanilino	fluoro	isopropyl	3.8 nM	11 nM
(R)-1-(hydroxymethyl)-isobutylamino	3-chloroanilino	bromo	isopropyl	43.8 nM	51 nM
(R)-1-(hydroxymethyl)-isobutylamino	3-chloroanilino	mercapto	isopropyl	83.7 nM	105.6 nM
(R)-1-(hydroxymethyl)-isobutylamino	3-chloroanilino	hydroxy	isopropyl	72.1 nM	89.5 nM
(R)-1-(hydroxymethyl)-isobutylamino	3-chloroanilino	amino	isopropyl	91.5 nM	92.8 nM
(R)-1-(hydroxymethyl)-isobutylamino	3-chloroanilino	2-hydroxy-ethylamino	isopropyl	1529.6 nM	1726.8 nM
(R)-1-(hydroxymethyl)-isobutylamino	3-chloroanilino	aminomethyl-amino	isopropyl	2521.3 nM	3287.2 nM

Table 4 shows the results of inhibitory activity of novel compounds against CDC2 and I κ B- α in comparison with the data on the prototype compounds. Most of the 2,6,8,9-tetrasubstituted purine derivatives showed marked inhibitory activity in *in vitro* kinase assays. Modification of 2,6,8-trisubstituted purines by small substituent in R8 position led usually to increase in cdk inhibitory activity of the tested compound.

EXAMPLE 13: Antimitotic activities of CDK inhibitors

Metaphase-arrested *Xenopus* egg extracts were prepared as described previously by Blow "J.Cell Biol." 122:993 (1993) and stored in liquid nitrogen. Demembranated *Xenopus* sperm nuclei were prepared as described by Blow & Laskey "Cell" 47:577 (1986). After thawing, extracts were supplemented with 25 mM phosphocreatine, 5 μ g/ml creatine phosphokinase, 250 μ g/ml cycloheximide, [α - 32 P]dATP (for DNA

synthesis assays). Demembranated sperm nuclei were added to a final sperm concentration of 3 ng/ μ l DNA extract and CDK inhibitor tested was then added at different concentrations. M-phase promoting factor inhibition by different CDK inhibitors was monitored 1.5 h after addition by assessing the amount of sperm nuclei that had been assembled into interphase nuclei, possessing a complete phase-dense nuclear envelope. DNA synthesis was assessed by releasing extract into interphase by the addition of 0.3 mM CaCl_2 and measuring the total amount of $[\alpha\text{-}^{32}\text{P}]\text{dATP}$ incorporation after 3 h by TCA co-precipitation.

At concentrations of CDK inhibitors (see Table 5) ranging from 0.1 - 2 μM , chromosomes remained highly condensed and no nuclear envelope was visible. At 4-6 μM and higher concentrations, interphase nuclei appeared with partially decondensed chromatin and an intact nuclear envelope. Replication was significantly inhibited at 1 - 5 μM CDK inhibitors tested. For the inhibition effect to become detectable, the first 15-min incubation of the interphase extract is probably sufficient.

Table 5. Antimitotic Activities of 2,6,8,9-Tetrasubstituted Purine Derivatives

SUBSTITUENT				Inhibition of MPF activity	Inhibition of DNA synthesis
C2	N6	C8	N9	IC_{50} (μM)	IC_{50} (μM)
hydroxyethylamino OLOMOUCINE	benzylamino	-	methyl	12	15
hydroxypropylamino	benzylamino	fluoro	isopropyl	2.6	3.2
(R)-1-(hydroxyethyl) propylamino	benzylamino	fluoro	isopropyl	1.5	2.4
(R)-1-(hydroxyethyl) propylamino	3-chloroanilino	fluoro	isopropyl	0.8	1.5
(R)-1-(hydroxyethyl) propylamino	3-chloro-4-carboxy- anilino	fluoro	isopropyl	0.1	0.25
(R)-1-(hydroxymethyl)- propylamino	3-chloroanilino	amino	isopropyl	3.12	3.3
(R)-1-(hydroxymethyl)- propylamino	3-chloroanilino	hydroxy	isopropyl	4.08	4.2
2-hydroxyethylamino	3-chloroanilino	2-hydroxyethylamino	isopropyl	41.7	63.5
(R)-1-(hydroxymethyl)-	3-chloroanilino	bromo	isopropyl	31.6	52.8

propylamino					
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EXAMPLE 14: *In vitro* Cytotoxic Activity of Novel Compounds

5 One of the parameters used, as the basis for colorimetric assays, is the metabolic activity of viable cells. For example, a microtiter assay, which uses the tetrazolium salt MTT, is now widely used to quantitate cell proliferation and cytotoxicity. For instance, this assay is used in drug screening programs and in chemosensitivity testing. Because only metabolically active cells cleave tetrazolium salts, these assays detect viable cells
10 exclusively. In the case of MTT assay, yellow soluble tetrazolium salt is reduced to coloured water-insoluble formazan salt. After it is solubilized, the formazan formed can easily and rapidly be quantified in a conventional ELISA plate reader at 570 nm (maximum absorbance). The quantity of reduced formazan corresponds to number of vital cells in the culture.

15 Human T-lymphoblastic leukemia cell line CEM; promyelocytic HL-60 and monocytic U937 leukemias; breast carcinoma cell lines MCF-7, BT549, MDA-MB-231; glioblastoma U87MG cells; cervical carcinoma cells HELA; sarcoma cells U2OS and Saos2; hepatocellular carcinoma HepG2; mouse fibroblasts NIH3T3; mouse immortalized bone marrow macrophages B2.4 and B10A.4; P388D1 and L1210
20 leukemia; B16 and B16F10 melanomas were used for routine screening of compounds. The cells were maintained in Nunc/Corning 80 cm² plastic tissue culture flasks and cultured in cell culture medium (DMEM with 5 g/l glucose, 2mM glutamine, 100 U/ml penicillin, 100 µg/ml streptomycin, 10% fetal calf serum and sodium bicarbonate).

 The cell suspensions that were prepared and diluted according to the particular
25 cell type and the expected target cell density (2.500-30.000 cells per well based on cell growth characteristics) were added by pipette (80µl) into 96/well microtiter plates. Inoculates were allowed a pre-incubation period of 24 hours at 37°C and 5% CO₂ for stabilisation. Four-fold dilutions of the intended test concentration were added at time zero in 20 µl aliquots to the microtiter plate wells. Usually, test compound was evaluated
30 at six 4-fold dilutions. In routine testing, the highest well concentration was 266.7 µM, but it can be the matter of change dependent on the agent. All drug concentrations were examined in duplicates. Incubations of cells with the test compounds lasted for 72 hours

at 37°C, in 5% CO₂ atmosphere and 100% humidity. At the end of incubation period, the cells were assayed by using the MTT. Ten microliters of the MTT stock solution were pipetted into each well and incubated further for 1-4 hours. After this incubation period, formazan was solubilized by addition of 100µl/well of 10% SDS in water (pH=5.5) followed by further incubation at 37°C overnight. The optical density (OD) was measured at 540nm with the Labsystem iEMS Reader MF (UK). The tumour cell survival (TCS) was calculated using the following equation: $TCS = (OD_{\text{drug exposed well}} / \text{mean } OD_{\text{control wells}}) \times 100\%$. The TCS₅₀ value, the drug concentration lethal to 50% of the tumour cells, was calculated from the obtained dose response curves.

Cytotoxicity of novel compounds was tested on panel of cell lines of different histogenetic and species origin (Table 6). We show here that equal activities were found in all tumour cell lines tested, however, the non-malignant cells, e.g. NIH3T3 fibroblasts and normal human lymphocytes, were resistant to synthetic CDK inhibitors induced cytotoxicity. As demonstrated in Table 6, IC₅₀ for NIH3T3 fibroblasts and normal human lymphocytes was always higher than 250 µM. Effective novel derivatives killed tumour cells in concentrations close to 1-5 µM. Notably, the identical effectiveness of purine derivatives was also found in cell lines bearing various mutations or deletions in cell cycle associated proteins, e.g. HL-60, BT549, Hela, U2OS, MDA-MB231, and Saos2 (Table 6). It indicates that these substances should be equally effective in tumours with various alterations of tumour suppresser genes, namely p53, Rb, etc. Importantly, this observation distinguishes novel compounds from flavopiridol and related compounds, as their biological activity is dependent on p53 status.

Table 6. Cytotoxicity of Novel Compounds for Different Cancer Cells

SUBSTITUENT				CEM	B16
C2	N6	C8	N9	IC ₅₀ (µM)	IC ₅₀ (µM)
2-hydroxyethylamino OLOMOUCINE	benzylamino	-	methyl	70	85.4
2-hydroxyethylamino	benzylamino	fluoro	methyl	45.2	62.8
2-hydroxyethylamino	benzylamino	bromo	methyl	165.2	172.8
2-hydroxyethylamino	benzylamino	mercapto	methyl	92.4	85.7

2-hydroxyethylamino	benzylamino	hydroxy	methyl	45.5	56.2
2-hydroxyethylamino	benzylamino	amino	methyl	84.2	67.1
2-hydroxyethylamino	benzylamino	2-hydroxyethylamino	methyl	>166.7	>166.7
2-hydroxyethylamino	benzylamino	aminomethylamino	methyl	114.3	126.5
3-hydroxypropylamino BOHEMINE	benzylamino	-	isopropyl	4.1	7.2
3-hydroxypropylamino	benzylamino	fluoro	isopropyl	3.93	5.5
3-hydroxypropylamino	benzylamino	bromo	isopropyl	116.9	76.4
3-hydroxypropylamino	benzylamino	methylthio	isopropyl	68.4	60.6
3-hydroxypropylamino	benzylamino	hydroxy	isopropyl	7.48	5.6
3-hydroxypropylamino	benzylamino	amino	isopropyl	8.52	6.8
3-hydroxypropylamino	benzylamino	2-hydroxyethylamino	isopropyl	>166.7	>166.7
3-hydroxypropylamino	benzylamino	2-aminoethylamino	isopropyl	29.9	2.5
(R)-1-(hydroxymethyl)- propylamino ROSCOVITINE	benzylamino		isopropyl	3.45	4.4
(R)-1-(hydroxymethyl)- propylamino	benzylamino	fluoro	isopropyl	3.38	3.2
(R)-1-(hydroxymethyl)- propylamino	benzylamino	bromo	isopropyl	48.4	59.2
(R)-1-(hydroxymethyl)- propylamino	benzylamino	mercapto	isopropyl	30.7	125.6
(R)-1-(hydroxymethyl)- propylamino	benzylamino	hydroxy	isopropyl	12.15	12.94
(R)-1-(hydroxymethyl)- propylamino	benzylamino	amino	isopropyl	12.17	12.85
(R)-1-(hydroxymethyl)- propylamino	benzylamino	2-hydroxyethylethoxy	isopropyl	140.2	124.95
(R)-1-(hydroxymethyl)- propylamino	benzylamino	1,2- dihydroxypropylamin o	isopropyl	128.2	147.5
(R)-1-(hydroxymethyl)- propylamino	benzylamino	2-aminoethylamino	isopropyl	96.1	>166.7
(R)-1-(hydroxymethyl)- propylamino	benzylamino	hydrazido	isopropyl	58	61,5
(R)-1-(hydroxymethyl)-	3-chloroanilino		isopropyl	2.4	3.5

isobutylamino PURVALANOL A					
(R)-1-(hydroxymethyl)- isobutylamino	3-chloroanilino	fluoro	isopropyl	2.5	3.5
(R)-1-(hydroxymethyl)- isobutylamino	3-chloroanilino	bromo	isopropyl	22.5	23.5
(R)-1-(hydroxymethyl)- isobutylamino	3-chloroanilino	mercapto	isopropyl	25.8	47.6
(R)-1-(hydroxymethyl)- isobutylamino	3-chloroanilino	hydroxy	isopropyl	11.8	11.3
(R)-1-(hydroxymethyl)- isobutylamino	3-chloroanilino	amino	isopropyl	11.5	11.8
(R)-1-(hydroxymethyl)- isobutylamino	3-chloroanilino	2-hydroxyethylamino	isopropyl	78	150
(R)-1-(hydroxymethyl)- isobutylamino	3-chloroanilino	2-aminomethylamino	isopropyl	56	130

EXAMPLE 15: Novel Compounds Induce Apoptosis in Tumour Cells.

To analyse the mechanisms of induced cytotoxicity by novel compounds, it is important to distinguish apoptosis from the other major form of cell death, necrosis. First, at the tissue level, apoptosis produces little or no inflammation, since the neighbouring cells, especially macrophages, rather than being released into the extracellular fluid, engulf shrunken portions of the cell. In contract, in necrosis, cellular contents are released into the extracellular fluid, and thus have an irritant affect on the nearby cells, causing inflammation. Second, at the cellular level, apoptotic cells exhibit shrinkage and blebbing of the cytoplasm, preservation of structure of cellular organelles including the mitochondria, condensation and margination of chromatin, fragmentation of nuclei, and formation of apoptotic bodies, though not all of these are seen in all cell types. Third, at the molecular level, a number of biochemical processes take an important role in induction of apoptosis. However, majority of them is not well understood, and they result in activation of proteases and nucleases, which finally destruct key biological macromolecules - proteins and DNA. For detection of apoptotic versus necrotic mode of cell death, two independent methods were employed:

assessment of morphology by electron microscopy and analysis of DNA fragmentation by flow-cytometry.

HL-60 cell line was cultured in 6-well culture plates with or without 70 μ M concentration of novel derivatives at 37°C and 5% CO₂ for 3-24 hours. Following the incubation, cells were pelleted, washed in Hank's buffered salt solution and processed as described below.

Cells were suspended in 2% glutaraldehyde/PBS, fixed overnight at 4°C, pelleted and embedded into 1% agar (Agar Noble, Difco) thereafter. Agar block containing fixed cells was epoxide polymerised, ultrathin sectioned, osmium tetroxide postfixed, uranium acetate contrasted and examined under electron microscope.

Initial phase contrast microscopy examinations indicated that treated HL-60 line exhibit typical morphological features of apoptotic cells, and it was later confirmed by electron microscopy. Typical morphological criteria of apoptosis were identified in cells treated with tetrasubstituted purine derivatives: chromatin condensation, nuclear fragmentation, cytoplasmatic blebbing, and formation of apoptotic bodies.

EXAMPLE 16: Immunosuppressive activity

One of the most important parameters of specific cellular immunity is proliferative response of lymphocytes to antigens or polyclonal mitogens. Majority of normal mammalian peripheral lymphocytes is resting cells. Antigens or nonspecific - polyclonal mitogens have capacity to activate lymphoid cells and it is accompanied with dramatic changes of intracellular metabolism (mitochondrial activity, protein synthesis, nucleic acids synthesis, formation of blastic cells and cellular proliferation). Compounds with ability to selectively inhibit lymphocyte proliferation are potent immunosuppressants. Variety of *in vitro* assays were developed to measure proliferative response of lymphocytes. The most commonly used is ³H-thymidine incorporation method.

During cell proliferation, the DNA has to be replicated before the cell is divided into two daughter cells. This close association between cell doublings and DNA synthesis is very attractive for assessing cell proliferation. If labeled DNA precursors are added to the cell culture, cells that are about to divide incorporate the labeled nucleotide into their DNA. Traditionally, those assays usually involve the use of radiolabeled

nucleosides, particularly tritiated thymidine ($[^3\text{H}]\text{-TdR}$). The amount of $[^3\text{H}]\text{-TdR}$ incorporated into the cellular DNA is quantified by liquid scintillation counting.

Human heparinized peripheral blood was obtained from healthy volunteers by cubital vein puncture. The blood was diluted in PBS (1:3) and mononuclear cells were
5 separated by centrifugation in Ficoll-Hypaque density gradient (Pharmacia, 1.077 g/ml) at 2200 g for 30 minutes. Following centrifugation, lymphocytes were washed in PBS and resuspended in cell culture medium (RMPI 1640, 2mM glutamine, 100 U/ml penicillin, 100 $\mu\text{g/ml}$ streptomycin, 10% fetal calf serum and sodium bicarbonate).

The cells were diluted at target density of 1.100.000 cells/ml were added by pipet
10 (180 μl) into 96/well microtiter plates. Four-fold dilutions of the intended test concentration were added at time zero in 20 μl aliquots to the microtiter plate wells. Usually, test compound was evaluated at six 4-fold dilutions. In routine testing, the highest well concentration was 266.7 μM . All drug concentrations were examined in duplicates. All wells with exception of unstimulated controls were activated with 50 μl
15 of concanavalin A (25 $\mu\text{g/ml}$). Incubations of cells with test compounds lasted for 72 hours at 37°C, in 5% CO_2 atmosphere and 100% humidity. At the end of incubation period, the cells were assayed by using the $[^3\text{H}]\text{-TdR}$:

Cell cultures were incubated with 0.5 μCi (20 μl of stock solution 500 $\mu\text{Ci/ml}$) per well for 6 hours at 37°C and 5% CO_2 . The automated cell harvester was used to lyse
20 cells in water and adsorb the DNA onto glass-fiber filters in the format of microtiter plate. The DNA incorporated $[^3\text{H}]\text{-TdR}$ was retained on the filter while unincorporated material passes through. The filters were dried at room temperature overnight, sealed into a sample bag with 10-12 ml of scintillant. The amount of $[^3\text{H}]\text{-TdR}$ present in each filter (in CCPM) was determined by scintillation counting in a Betaplate liquid
25 scintillation counter. The effective dose of immunosuppressant (ED) was calculated using the following equation: $\text{ED} = (\text{CCPM}_{\text{drug exposed well}} / \text{mean CCPM}_{\text{control wells}}) \times 100\%$. The ED_{50} value, the drug concentration inhibiting proliferation of 50% of lymphocytes, was calculated from the obtained dose response curves.

To evaluate immunosuppressive activity of tetrasubstituted adenines, their ability
30 to inhibit polyclonal mitogen induced proliferation of normal human lymphocytes was analyzed (Table 7). Our data demonstrate that these compounds have only marginal activity on ^3H -thymidine incorporation, nonetheless, they efficiently inhibit proliferation

of activated lymphocytes. Effective immunosuppressive dose of tetrasubstituted derivatives under in vitro conditions was close to 1-20 μM .

Table 7 Immunosuppressive effects of compounds on spontaneous and mitogen
5 activated proliferation of lymphocytes

SUBSTITUENT				spontane- ous	mitogen activated
C2	N6	C8	N9	IC ₅₀ (μM)	IC ₅₀ (μM)
hydroxyethylamino OLOMOUCIN	benzylamino	-	methyl	245	72.1
2-hydroxyethylamino	benzylamino	bromo	methyl	228	68.3
2-hydroxyethylamino	benzylamino	mercapto	methyl	250	83.5
2-hydroxyethylamino	benzylamino	hydroxy	methyl	201	48.6
2-hydroxyethylamino	benzylamino	amino	methyl	170.3	26.7
2-hydroxyethylamino	benzylamino	2-hydroxy- ethylamino	methyl	250	169.3
2-hydroxyethylamino	benzylamino	aminomethyl- amino	methyl	250	113.5
3-hydroxypropylamino BOHEMIN	benzylamino	-	isopropyl	172	5.7
3-hydroxypropylamino	benzylamino	bromo	isopropyl	178	5.9
3-hydroxypropylamino	benzylamino	mercapto	isopropyl	197	18.4
3-hydroxypropylamino	benzylamino	hydroxy	isopropyl	163	2.1
3-hydroxypropylamino	benzylamino	amino	isopropyl	143	1.59
3-hydroxypropylamino	benzylamino	2-hydroxy- ethylamino	isopropyl	181	29.3
3-hydroxypropylamino	benzylamino	aminomethyl- amino	isopropyl	174	26.8
(R)-1-(hydroxymethyl)- propylamino ROSCOVITIN	benzylamino		isopropyl	164	6.4
(R)-1-(hydroxymethyl)- propylamino	benzylamino	bromo	isopropyl	168	17.1
(R)-1-(hydroxymethyl)- propylamino	benzylamino	mercapto	isopropyl	174	18.5

(R)-1-(hydroxymethyl)-propylamino	benzylamino	hydroxy	isopropyl	153	12.8
(R)-1-(hydroxymethyl)-propylamino	benzylamino	amino	isopropyl	149	13.3
(R)-1-(hydroxymethyl)-propylamino	benzylamino	2-hydroxyethyl-amino	isopropyl	182	31.5
(R)-1-(hydroxymethyl)-propylamino	benzylamino	aminomethyl-amino	isopropyl	177	25.8
(R)-1-(hydroxymethyl)-isobutylamino PURVALANOL A	3-chloroanilino		isopropyl	158	5.7
(R)-1-(hydroxymethyl)-isobutylamino	3-chloroanilino	bromo	isopropyl	142	13.7
(R)-1-(hydroxymethyl)-isobutylamino	3-chloroanilino	mercapto	isopropyl	164	26.9
(R)-1-(hydroxymethyl)-isobutylamino	3-chloroanilino	hydroxy	isopropyl	126	2.98
(R)-1-(hydroxymethyl)-isobutylamino	3-chloroanilino	amino	isopropyl	103	3.1
(R)-1-(hydroxymethyl)-isobutylamino	3-chloroanilino	2-hydroxyethyl-amino	isopropyl	205	150
(R)-1-(hydroxymethyl)-isobutylamino	3-chloroanilino	aminomethyl-amino	isopropyl	230	138

EXAMPLE 17: Antiviral Activity

The activity of the compounds against HIV-1 and HIV-2 induced cytopathicity was examined in human lymphocyte MT-4 cells. The cells (300 000 cells/ml) were infected with 100 CCID₅₀ (1 CCID₅₀ is a virus quantity which causes cytopathicity effect in 50% of the cells under the experimental conditions) of HIV-1 or HIV-2 and added to 200 µl wells of a microtiter plate containing different dilutions of the tested compounds. The infected cell cultures were incubated at 37°C for 5 days in a humidified CO₂ incubator. The cytopathicity of the virus was examined by determination of MT-4 cell viability by trypan blue dye staining. The results are summarised in Table 8 with comparison on the prototype compounds.

Table 8 also shows the results of activity testing of novel compounds against MSV-induced transformation in murine embryo fibroblast C3H/3T3 cells. The cells were seeded in 1ml wells of 48-well plates and exposed to 80 PFU (plaque forming units) for 60 - 90 min. The virus was subsequently removed and culture medium
 5 containing appropriate concentrations of the tested compounds was added (1 ml per well). At day 6-post infection, MSV-induced transformation of the cell culture was examined microscopically.

Table 8. Antiretroviral Activity of Purine Compounds, wherein R9 is either PMP (2-phosphonomethoxypropyl)group or PME (2-phosphonomethoxyethyl)group. ($\mu\text{g/ml}$),
 10 (R2 = NH₂).

R6	R8	MSV	HIV-1		HIV-2	
PME-derivatives			MT-4	CEM	MT-4	CEM
amino		0,6	2,67	6.9	ND	ND
amino	fluoro	0,57	3,12	7.2	ND	ND
amino	mercapto	1.23	5.87	10.8	ND	ND
amino	hydroxy	0.42	1.95	5.2	ND	ND
amino	amino	0.58	1.98	4.8	ND	ND
amino	2-hydroxyethylamino	1.54	8.33	10.8	ND	ND
amino	aminomethylamino	1.32	7.54	9.3	ND	ND
cyclohexylamino		0.26	5.7	>20	4.8	>20
cyclohexylamino	fluoro	0.24	6.3	>20	4.6	>20
cyclohexylamino	mercapto	0.35	8.9	>20	8.5	>20
cyclohexylamino	hydroxy	0.21	4.3	>20	2.1	>20
cyclohexylamino	amino	0.19	3.5	>20	1.6	>20
cyclohexylamino	2-hydroxyethylamino	0.45	12.7	>20	12.7	>20
cyclohexylamino	aminomethylamino	0.43	10.8	>20	9.5	>20
benzylamino		1.5	50	>20	49	>20
benzylamino	fluoro	1.3	47	>20	45	>20
benzylamino	mercapto	1.8	56	>20	57	>20
benzylamino	hydroxy	0.9	45	>20	32	>20
benzylamino	amino	0.8	48	>20	31	>20
benzylamino	2-hydroxyethylamino	1.7	67	>20	48	>20
benzylamino	aminomethylamino	1.4	55	>20	52	>20

R6	R8	MSV	HIV-1		HIV-2	
PMP-derivatives			MT-4	CEM	MT-4	CEM
amino		0,07	0,29	10	0.24	10
amino	fluoro	0.06	3,12	ND	3.54	ND
amino	mercapto	0.15	4.18	ND	4.15	ND
amino	hydroxy	0.05	0.25	ND	0.21	ND
amino	amino	0.06	0.19	ND	0.20	ND
amino	2-hydroxyethylamino	0.35	5.16	ND	4.87	ND
amino	aminomethylamino	0.42	4.58	ND	4.65	ND
cyclohexylamino		3.78	3.4	4.5	5.8	8.5
cyclohexylamino	fluoro	2,54	3.2	4.1	4.6	8.3
cyclohexylamino	mercapto	6.32	10.1	>20	11.2	>20
cyclohexylamino	hydroxy	1.37	2.1	5.2	3.2	7.8
cyclohexylamino	amino	1.25	1.8	4.7	2.8	8.1
cyclohexylamino	2-hydroxyethylamino	5.42	12.7	>20	25.1	>20
cyclohexylamino	aminomethylamino	4.98	9.8	>20	18.6	>20
benzylamino		0.3	10.3	11.6	8.3	12.5
benzylamino	fluoro	0.27	8.5	12.5	6.4	11.1
benzylamino	mercapto	1.35	21.8	>20	>20	>20
benzylamino	hydroxy	0.21	3.7	10.3	4.3	11.1
benzylamino	amino	0.18	2.9	10.7	3.8	11.9
benzylamino	2-hydroxyethylamino	2.17	19.5	>20	>20	>20
benzylamino	aminomethylamino	1.89	16.8	>20	>20	>20

Most of the PMP (9-(2-phosphonomethoxypropyl)derivative) and PME (9-(2-phosphonomethoxyethyl)derivative) compounds of the formula I showed marked anti-HIV activity *in vitro*. HIV-1 and HIV-2 did not differ in their sensitivity to the test compounds. (R)-PMP compounds were markedly inhibitory to retroviruses at 2 - 3 $\mu\text{g/ml}$ and not toxic to the cells at 100 $\mu\text{g/ml}$. Its selectivity index (ratio cytotoxic dose/antivirally active dose) proved superior over that of the prototype compound PME. The (S)-enantiomer of PME was devoid of marked antiretroviral activity. (R)-PMPD were exquisitely inhibitory to retrovirus replication (EC_{50} 0.01 -0.1 $\mu\text{g/ml}$) and not toxic to the cells at 100 $\mu\text{g/ml}$. It proved superior over PMEA and other prototype compounds

in terms of both antiviral activity and lack of toxicity. Its selectivity index was higher than 2000 for HIV-1 and HIV-2.

EXAMPLE 18: Dry Capsules

- 5 5000 capsules, each of which contain 0.25 g of one of the compounds of the formula I mentioned in the preceding Examples as active ingredient, are prepared as follows:

Composition

	Active ingredient	1250 g
	Talc	180 g
10	Wheat starch	120 g
	Magnesium stearate	80 g
	Lactose	20 g

- Preparation process: The powdered substances mentioned are pressed through a sieve of mesh width 0.6 mm. Portions of 0.33 g of the mixture are transferred to gelatine capsules
15 with the aid of a capsule-filling machine.

EXAMPLE 19: Soft Capsules

- 5000 soft gelatine capsules, each of which contain 0.05 g of one of the compounds of the formula I mentioned in the preceding Examples as active ingredient, are prepared as
20 follows:

Composition

	Active ingredient	250 g
	Lauroglycol	2 litres

- Preparation process: The powdered active ingredient is suspended in Lauroglykol®
25 (propylene glycol laurate, Gattefossé S.A., Saint Priest, France) and ground in a wet-pulveriser to a particle size of about 1 to 3 µm. Portions of in each case 0.419 g of the mixture are then transferred to soft gelatine capsules by means of a capsule-filling machine.

EXAMPLE 20: Soft Capsules

5000 soft gelatine capsules, each of which contain 0.05 g of one of the compounds of the formula I mentioned in the preceding or following Examples as active ingredient, are prepared as follows:

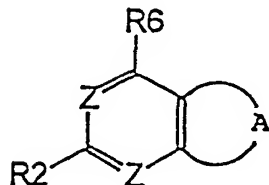
5 Composition

Active ingredient	250 g	-
PEG 400	1 litre	
Tween 80	1 litre	

10 Preparation process: The powdered active ingredient is suspended in PEG 400 (polyethylene glycol of Mr between 380 and about 420, Sigma, Fluka, Aldrich, USA) and Tween[®] 80 (polyoxyethylene sorbitan monolaurate, Atlas Chem. Inc., USA, supplied by Sigma, Fluka, Aldrich) and ground in a wet-pulveriser to a particle size of about 1 to 3 mm. Portions of in each case 0.43 g of the mixture are then transferred to soft gelatine capsules by means of a capsule-filling machine.

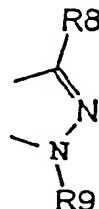
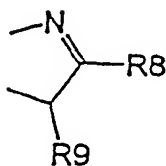
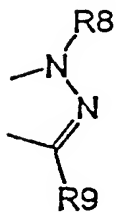
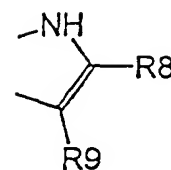
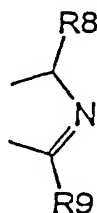
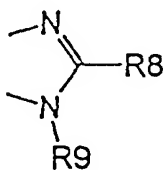
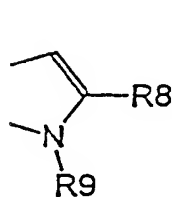
CLAIMS

1. Substituted nitrogen heterocyclic derivatives of the formula I wherein



(I)

10 A is a divalent group selected from the ensemble consisting of



Z is N or CH, provided that at least one Z being N;

R2 and R6 are independent of one another, represent H, halogen, alkyl, substituted alkyl, cycloalkyl, substituted cycloalkyl, cycloalkyl alkyl, arylalkyl, heteroalkyl, heteroarylalkyl, cycloheteroalkyl alkyl or R6'-X wherein

25 X is an -NH-, -N(C₁-C₆-alkyl)-, -O- or -S- moiety;

R6' is H, alkyl, substituted alkyl, acyl, amido, sulfo, cycloalkyl, substituted cycloalkyl, aryl, substituted aryl, heterocycle, heteroaryl, substituted heteroaryl, arylalkyl, cycloheteroalkyl, substituted cycloheteroalkyl, heteroarylalkyl, heteroalkyl, cycloalkyl alkyl and cycloheteroalkyl alkyl;

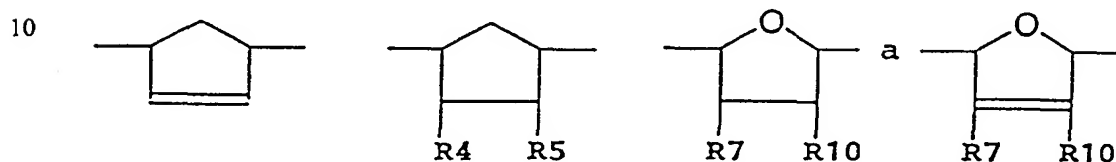
30 R8 is halogen, hydroxyl, amino, carboxyl, cyano, nitro, amido, sulfo, sulfamido, carbamino, alkyl, substituted alkyl, cycloalkyl, substituted cycloalkyl, arylalkyl, heteroalkyl, heteroarylalkyl, cycloalkyl alkyl, cycloheteroalkyl alkyl or R8'-X wherein

X is -NH-, -N(alkyl)-, -O- or -S- moiety and

R8' is according to any one of the substituents defined above for R2' or R6'.

R9 is alkyl, substituted alkyl, acyl, carboxyl, amido, sulfamino, cycloalkyl, substituted cycloalkyl, cycloalkyl alkyl, heteroalkylcycloalkyl alkyl, heterocycloalkyl, substituted heterocycloalkyl, aryl, substituted aryl, heterocycle, heteroaryl, substituted heteroaryl, arylalkyl, heteroarylalkyl, heteroalkyl or -B-R9'- wherein

B is -CH₂-, -(CH₂)₂-, -CH(CH₃)CH₂-, -CH(CH₂F)CH₂-, -CH(CH₂OH)CH₂-, or the groups of the following structure,



wherein the left hand bond is linked to nitrogen of 5-membered ring of compounds of the formula I;

R4 and R5, that are independent of one another, represent hydrogen, hydroxyl, halogen, amino, acyloxy substituent having 1-5 carbon atoms, alkoxy substituent having 1-5 carbon atoms, alkylmercapto substituent having 1-5 carbon atoms, alkylamino substituent having 1-5 carbon atoms and dialkylamino in which each alkyl substituent has 1-5 carbon atoms;

R7 and R10, that are independent of one another, represent H or alkyl substituent having 1-10 carbon atoms;

R9 is -(CH₂)_n-R9', wherein n = 1-2 and the

R9' is -X(CH₂)_mY wherein

25 X is -O-, -S-, -NH- or -N (alkyl)- substituent having 1-6 carbon atoms;

m = 1-2;

Y is carboxyl, amido, sulfo, sulfamino, hydroxyl, carboxyl, mercapto, carbylmercapto, amino, alkylamino, carbamino -PO(OH)₂, -PO(O-C₁-C₆-alkyl)₂, -PO(NH-C₁-C₆-alkyl)₂, PO(O-C₁-C₆-alkyl)(NH-C₁-C₆-alkyl), -PO(OH)(O-C₁-C₆-alkyl);

30 -PO(OH)(NH-C₁-C₆-alkyl) or

-(CH₂CHD)-R^{9'}, wherein

R^{9'}, X, m and Y are as defined above and

D is alkyl, substituted alkyl, -PO(OH)₂, -PO(OH)(O-C₁-C₆-alkyl), -PO(OH)(NH-C₁-C₆-alkyl).

5

The above not yet defined generic group having meanings as introduced in the following legend:

“halogen“

refers to fluorine, bromine, chlorine, and iodine atoms;

10 “alkyl“

refers to

- a branched or unbranched alkyl group having 1-6 carbon atoms,
- a branched or unbranched alkenyl group having 1-6 carbon atoms,
- a branched or unbranched alkynyl group having 1-6 carbon atoms;

15 “substituted alkyl“

refers to a branched or unbranched alkyl, alkenyl or alkynyl group having 1-6 carbon atoms and having substituted by one or more substituents selected from the group consisting of hydroxyl, mercapto, carbylmercapto, halogen, carbyloxy, amino, amido, carboxyl, cycloalkyl, sulfo or acyl. These substituent generic groups having the meanings being identical with the definitions of the corresponding groups as defined in this legend;

20

“carbyloxy“

denotes the group -OR_a, where R_a is alkyl, substituted alkyl, aryl, substituted aryl, arylalkyl, substituted arylalkyl, cycloalkyl, substituted cycloalkyl, heterocycloalkyl or substituted heterocycloalkyl whereas these generic groups have meanings being identical with definitions of the corresponding groups as defined in this legend;

25

“carbylmercapto“

denotes the group -SR_b, where R_b is alkyl, substituted alkyl, aryl, substituted aryl, arylalkyl, substituted arylalkyl, cycloalkyl, substituted cycloalkyl, heterocycloalkyl or substituted heterocycloalkyl whereas these generic groups have meanings being identical with definitions of the corresponding groups as defined in this legend;

30

“sulfo“

denotes the group $-\text{SO}_3\text{R}_c$, where R_c is

- hydrogen,
- a branched or unbranched alkyl group having 1-6 carbon atoms,
- a branched or unbranched alkenyl group having 1-6 carbon atoms,
- 5 • a branched or unbranched alkynyl group having 1-6 carbon atoms,

a branched or unbranched alkyl, alkenyl or alkynyl group having 1-6 carbon atoms and being substituted by one or more substituents selected from the group consisting of hydroxyl, mercapto, carbylmercapto, halogen, carbyloxy, amino, amido, carboxyl, cycloalkyl, sulfo or acyl, whereas these generic groups have meanings being identical with the definitions of the corresponding groups as defined in this legend;

“sulfamino“

denotes the group $-\text{NHSO}_3\text{R}_d$, wherein R_d is

- hydrogen,
- a branched or unbranched alkyl group having 1-6 carbon atoms,
- 15 • a branched or unbranched alkenyl group having 1-6 carbon atoms,
- a branched or unbranched alkynyl group having 1-6 carbon atoms,

a branched or unbranched alkyl, alkenyl or alkynyl group having 1-6 carbon atoms and being substituted by one or more substituents selected from the group consisting of hydroxyl, mercapto, carbylmercapto, halogen, carbyloxy, amino, amido, carboxyl, cycloalkyl, sulfo or acyl, whereas these generic substituents have meanings being identical with definitions of the corresponding groups as defined in this legend;

“acyl“

denotes the group $-\text{C}(\text{O})\text{R}_e$, where R_e is hydrogen, alkyl, substituted alkyl, aryl, substituted aryl, arylalkyl, substituted arylalkyl, cycloalkyl, substituted cycloalkyl, whereas these generic groups have meanings being identical with definitions of the corresponding groups as defined in this legend;

“aryloxy“

denotes the group $-\text{OAr}$, where Ar is an aryl, substituted aryl, heteroaryl or substituted heteroaryl whereas these generic groups have meanings being identical with definitions of the corresponding groups as defined in this legend;

“alkylamino“

denotes the group $-NR_fR_g$, where R_f and R_g , that are independent of one another, represent hydrogen, alkyl, substituted alkyl, aryl, substituted aryl, heteroaryl or substituted heteroaryl, whereas these generic substituents have meanings being identical with definitions of the corresponding groups defined herein;

- 5 “amido“ denotes the group $-C(O)NR_hR_i'$, where R_h and R_i' may independently be hydrogen, alkyl, substituted alkyl, aryl, substituted aryl, heteroaryl, substituted heteroaryl whereas these generic groups have meanings being identical with definitions of the corresponding groups as defined in this legend;

“carboxyl“

- 10 denotes the group $-C(O)OR_j$, where R_j is hydrogen, alkyl, substituted alkyl, aryl, substituted aryl, heteroaryl or substituted heteroaryl, whereas these generic substituents have meanings being identical with definitions of the corresponding groups defined herein;

“carbamino“

- 15 denotes the group $-NHCOR_k$, where R_k may be hydrogen, alkyl, substituted alkyl, heterocycle, aryl, substituted aryl, heteroaryl and substituted heteroaryl whereas these generic groups have meanings being identical with definitions of the corresponding groups as defined in this legend;

“aryl“

- 20 refers to an aromatic carbocyclic group having from 1 to 18 carbon atoms and being composed of at least one aromatic or multiple condensed rings in which at least one of which being aromatic;

“substituted aryl“

- refers to an aromatic carbocyclic group having from 1 to 18 carbon atoms and being
25 composed of at least one aromatic ring or of multiple condensed rings at least one of which being aromatic. The ring(s) are optionally substituted with one or more substituents selected from the group consisting of halogen, alkyl, hydroxyl, carbylmercapto, alkylamino, carbyloxy, amino, amido, carboxyl, nitro, mercapto or sulfo, whereas these generic substituent group have meanings being identical with
30 definitions of the corresponding groups as defined in this legend;

“heterocycle“

refers to a heterocyclic group having from 4 to 9 carbon atoms and at least one heteroatom selected from the group consisting of N, O or S;

“heteroaryl“

refers to a heterocyclic group having from 4 to 9 carbon atoms and at least one heteroatom selected from the group consisting of N, O or S with at least one ring of this group being aromatic;

“substituted heteroaryl“

refers to a heterocyclic group having from 4 to 9 carbon atoms and at least one heteroatom selected from the group consisting of N, O or S with at least one ring of this group being aromatic and this group being substituted with one or more substituents selected from the group consisting of halogen, alkyl, carbyloxy, carbylmercapto, alkylamino, amido, carboxyl, hydroxyl, nitro, mercapto or sulfo, whereas these generic substituent group have meanings being identical with definitions of the corresponding groups as defined in this legend;

“arylalkyl“

denotes the group $-R_i-Ar$ where R_i is

- a branched or unbranched alkyl group having 1-6 carbon atoms,
- a branched or unbranched alkenyl group having 1-6 carbon atoms,
- a branched or unbranched alkynyl group having 1-6 carbon atoms,

and Ar is

an aromatic carbocyclic group having from 1 to 18 carbon atoms and being composed of at least one aromatic ring or of multiple condensed rings at least one of which being aromatic and the group being optionally substituted with one or more substituents selected from the group consisting of halogen, alkyl, hydroxyl, carbylmercapto, alkylamino, carbyloxy, amino, amido, carboxyl, nitro, mercapto or sulfo, whereas these generic substituent group have meanings being identical with definitions of the corresponding groups as defined in this legend;

“heteroalkyl“

denotes the group $-R_m-L$ where L is

- a branched or unbranched alkyl group having 1-6 carbon atoms,
- a branched or unbranched alkenyl group having 1-6 carbon atoms,
- a branched or unbranched alkynyl group having 1-6 carbon atoms,

- a branched or unbranched alkyl, alkenyl or alkynyl group having 1-6 carbon atoms and being substituted by one or more substituents selected from the group consisting of hydroxyl, mercapto, carbylmercapto, halogen, carbyloxy, amino, amido, carboxyl, cycloalkyl, sulfo or acyl, whereas these generic substituent group have meanings being identical with definitions of the corresponding groups as defined in this legend and L is

a heterocyclic group having from 4 to 9 carbon atoms and at least one heteroatom selected from the group consisting of N, O or S and the group being unsubstituted or substituted with one or more substituents selected from the group consisting of halogen, alkyl, alkoxy, alkylmercapto, alkylamino, amido, carboxyl, hydroxy, nitro, mercapto or sulfo, whereas these generic substituents have meanings being identical with definitions of the corresponding groups as defined in this legend;

“heteroarylalkyl”

denotes the group -R_n-G- where R_n is

- a branched or unbranched alkyl group having 1-6 carbon atoms,
- a branched or unbranched alkenyl group having 1-6 carbon atoms,
- a branched or unbranched alkynyl group having 1-6 carbon atoms,
- a branched or unbranched alkyl, alkenyl or alkynyl group having 1-6 carbon atoms and being substituted by one or more substituents selected from the group consisting of hydroxyl, mercapto, carbylmercapto, halogen, carbyloxy, amino, amido, carboxyl, cycloalkyl, sulfo or acyl, whereas these generic substituent group have meanings being identical with definitions of the corresponding groups as defined in this legend and G is

a heterocyclic group having from 4 to 9 carbon atoms and at least one heteroatom selected from the group consisting of N, O or S with at least one ring of which being aromatic and the group being optionally substituted with one or more substituents selected from the group consisting of halogen, alkyl, carbyloxy, carbylmercapto, alkylamino, amido, carboxyl, hydroxyl, nitro, mercapto or sulfo, whereas these generic substituent group have meanings being identical with definitions of the corresponding groups as defined in this legend;

“cycloalkyl”

refers to a monocyclic or polycyclic alkyl group containing 3 to 15 carbon atoms;

“substituted cycloalkyl”

refers to a monocyclic or polycyclic alkyl group containing 3 to 15 carbon atoms and being substituted by one or more substituents selected from the group consisting of halogen, alkyl, substituted alkyl, carbyloxy, carbylmercapto, aryl, nitro, mercapto or sulfo, whereas these generic substituent groups have meanings being identical with definitions of the corresponding groups as defined in this legend;

“heterocycloalkyl”

refers to a monocyclic or polycyclic alkyl group containing 3 to 15 carbon atoms which at least one ring carbon atom of its cyclic structure being replaced with a heteroatom selected from the group consisting of N, O, S or P;

“substituted heterocycloalkyl”

refers to a monocyclic or polycyclic alkyl group containing 3 to 15 carbon atoms which at least one ring carbon atom of its cyclic structure being replaced with a heteroatom selected from the group consisting of N, O, S or P and the group is containing one or more substituents selected from the group consisting of halogen, alkyl, substituted alkyl, carbyloxy, carbylmercapto, aryl, nitro, mercapto or sulfo, whereas these generic substituent group have meanings being identical with definitions of the corresponding groups as defined in this legend;

“cycloalkyl alkyl”

denotes the group $-R_0-J$ where R_0 is

- a branched or unbranched alkyl group having 1-6 carbon atoms,
- a branched or unbranched alkenyl group having 1-6 carbon atoms,
- a branched or unbranched alkynyl group having 1-6 carbon atoms,
- a branched or unbranched alkyl, alkenyl or alkynyl group having 1-6 carbon atoms and being substituted by one or more substituents selected from the group consisting of hydroxyl, mercapto, carbylmercapto, halogen, carbyloxy, amino, amido, carboxyl, cycloalkyl, sulfo or acyl, whereas these generic substituent group have meanings being identical with definitions of the corresponding groups as defined in this legend

and J is

- a monocyclic or polycyclic alkyl group containing 3 to 15 carbon atoms;
- a monocyclic or polycyclic alkyl group containing 3 to 15 carbon atoms which contains one or more substituents selected from the group consisting of halogen,

alkyl, substituted alkyl, carbyloxy, carbylmercapto, aryl, nitro, mercapto or sulfo, whereas these generic substituent groups have meanings being identical with definitions of the corresponding groups as defined in this legend and

“heterocycloalkylalkyl”

5 denotes the group $-R_pV$ where R_p is

- a branched or unbranched alkyl group having 1-6 carbon atoms,
 - a branched or unbranched alkenyl group having 1-6 carbon atoms,
 - a branched or unbranched alkynyl group having 1-6 carbon atoms,
 - a branched or unbranched alkyl, alkenyl or alkynyl group having 1-6 carbon atoms
- 10 and being substituted by one or more substituents selected from the group consisting of hydroxyl, mercapto, carbylmercapto, halogen, carbyloxy, amino, amido, carboxyl, cycloalkyl, sulfo or acyl, whereas these generic substituent groups have meanings being identical with definitions of the corresponding groups as defined in this legend

and V is

- 15 • a monocyclic or polycyclic alkyl group containing 3 to 15 carbon atoms with at least one being replaced with a heteroatom selected from the group consisting of N, O, S or P;
 - a monocyclic or polycyclic alkyl group containing 3 to 15 carbon atoms with at least one being replaced with a heteroatom selected from the group consisting of N, O, S
- 20 or P and the group contains one or more substituents selected from the group consisting of halogen, alkyl, substituted alkyl, carbyloxy, carbylmercapto, aryl, nitro, mercapto or sulfo, whereas these generic substituent groups have meanings being identical with definitions of the corresponding groups as defined in this legend and the pharmaceutically acceptable acid salts, racemates and optical isomers thereof.

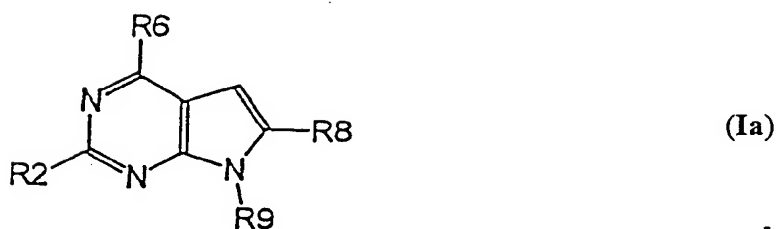
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2. Substituted nitrogen heterocyclic derivatives of the formula I according to claim 1, wherein $R_6 = H$ and R_2 , R_8 and R_9 are as defined in claim 1.

3. Substituted nitrogen heterocyclic derivatives of the formula I according to claim 1, wherein $R_2 = H$ and R_6 , R_8 and R_9 are as defined in claim 1.

30

4. Substituted nitrogen heterocyclic derivatives according to claim 1 of the formula Ia,



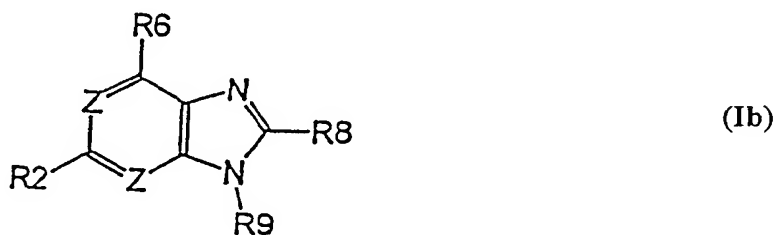
wherein R2, R6, R8 and R9 are as defined in claim 1.

10 5. Substituted nitrogen heterocyclic derivatives of the formula **Ia** according to claim 4, wherein R6 = H and R2, R8 and R9 are as defined in claim 1.

6. Substituted nitrogen heterocyclic derivatives of the formula **Ia** according to claim 4, wherein R2 = H and R6, R8 and R9 are as defined in claim 1.

15

7. Substituted nitrogen heterocyclic derivatives according to claim 1 of the formula **Ib**,



wherein R2, R6, R8 and R9 are as defined above in claim 1.

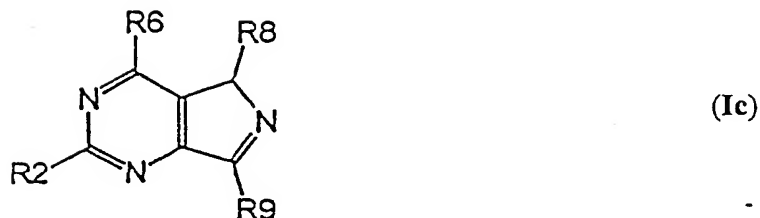
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8. Substituted nitrogen heterocyclic derivatives of the formula **Ib** according to claim 7, wherein R6 = H and R2, R8 and R9 are as defined in claim 1.

9. Substituted nitrogen heterocyclic derivatives of the formula **Ib** according to claim 7, wherein R2 = H and R6, R8 and R9 are as defined in claim 1.

30

10. Substituted nitrogen heterocyclic derivatives according to claim 1 of the formula **Ic**,

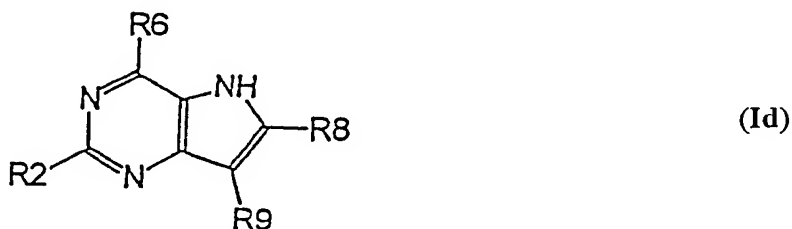


wherein R2 , R6, R8 and R9 are as defined above in claim 1.

10 11. Substituted nitrogen heterocyclic derivatives of the formula **Ic** according to claim 10, wherein R6 = H and R2, R8 and R9 are as defined in claim 1.

12. Substituted nitrogen heterocyclic derivatives of the formula **Ic** according to claim 10, wherein R2 = H and R6, R8 and R9 are as defined in claim 1.

15 13. Substituted nitrogen heterocyclic derivatives according to claim 1 of the formula **Id**,



wherein R2, R6, R8 and R9 are as defined above in claim 1.

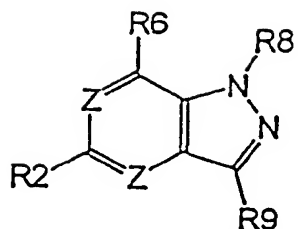
25 14. Substituted nitrogen heterocyclic derivatives of the formula **Id** according to claim 13, wherein R6 = H and R2, R8 and R9 are as defined in claim 1.

15. Substituted nitrogen heterocyclic derivatives of the formula **Id** according to claim 13, wherein R2 = H and R6, R8 and R9 are as defined in claim 1.

30

16. Substituted nitrogen heterocyclic derivatives according to claim 1 of the formula **Ie**,

(Ie)

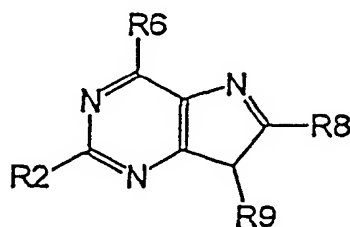


wherein R₂, R₆, R₈ and R₉ are as defined in claim 1.

17. Substituted nitrogen heterocyclic derivatives of the formula **Ie** according to claim 16, wherein R₆ = H and R₂, R₈ and R₉ are as defined in claim 1.

18. Substituted nitrogen heterocyclic derivatives of the formula **Ie** according to claim 16, wherein R₂ = H and R₆, R₈ and R₉ are as defined in claim 1.

19. Substituted nitrogen heterocyclic derivatives according to claim 1 of the formula **If**,



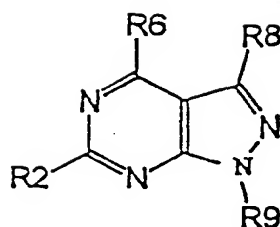
(If)

wherein R₂, R₆, R₈ and R₉ are as defined in claim 1.

20. Substituted nitrogen heterocyclic derivatives of the formula **If** according to claim 19, wherein R₆ = H and R₂, R₈ and R₉ are as defined in claim 1.

21. Substituted nitrogen heterocyclic derivatives of the formula **If** according to claim 19, wherein R₂ = H and R₆, R₈ and R₉ are as defined in claim 1.

22. Substituted nitrogen heterocyclic derivatives according to claim 1 of the formula **Ig**,



(Ig)

wherein R2, R6, R8 and R9 are as defined in claim 1.

23. Substituted nitrogen heterocyclic derivatives of the formula Ig according to claim 22, wherein R6 = H and R2, R8 and R9 are as defined in claim 1.

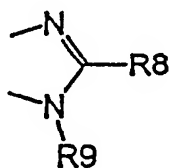
24. Substituted nitrogen heterocyclic derivatives of the formula Ig according to claim 22, wherein R2 = H and R6, R8 and R9 are as defined in claim 1.

25. Substituted nitrogen heterocyclic derivatives of the formula I according to claim 1 selected from the group consisting of 2-(1-hydroxymethylpropylamino)-6-benzylamino-8-chloro (or hydroxy, bromo, fluoro, amino, amido, carboxy, cyano, methylamino, thio, methylthio, ω -hydroxyalkylamino, ω -hydroxyalkyloxy, ω -carboxyalkylamino, ω -aminoalkylamino, ω -fosfonoalkylamino, ω -fosfonoalkyloxy, propinyl)-9-isopropylpurine, 2-(2-aminopropylamino)-6-benzylamino-8-chloro (or hydroxy, bromo, fluoro, amino, amido, carboxy, cyano, methylamino, thio, methylthio, ω -hydroxyalkylamino, ω -hydroxyalkyloxy, ω -carboxyalkylamino, ω -aminoalkylamino, ω -fosfonoalkylamino, ω -fosfonoalkyloxy, propinyl)-9-isopropylpurine, 2-(2-hydroxypropylamino)-6-benzylamino-8-chloro (or hydroxy, bromo, fluoro, amino, amido, carboxy, cyano, methylamino, thio, methylthio, ω -hydroxyalkylamino, ω -hydroxyalkyloxy, ω -carboxyalkylamino, ω -aminoalkylamino, ω -fosfonoalkylamino, ω -fosfonoalkyloxy, propinyl)-9-isopropylpurine, 2-diethylamino-6-(4-methoxybenzylamino)-8-chloro (or hydroxy, bromo, fluoro, amino, amido, carboxy, cyano, methylamino, thio, methylthio, ω -hydroxyalkylamino, ω -hydroxyalkyloxy, ω -carboxyalkylamino, ω -aminoalkylamino, ω -fosfonoalkylamino, ω -fosfonoalkyloxy, propinyl)-9-isopropylpurine, 2-(2-hydroxypropylamino)-6-(3-chloroanilino)-8-chloro (or hydroxy, bromo, amino, C₁-C₆ alkyl, methyl, ethyl, propyl, isopropyl, vinyl, allyl, propargyl)-9-isopropylpurine, 2-(2-hydroxypropylamino)-6-(3-

chloro-4-carboxyanilino)-8-chloro (or hydroxy, bromo, fluoro, amino, amido, carboxy, cyano, methylamino, thio, methylthio, ω -hydroxyalkylamino, ω -hydroxyalkyloxy, ω -carboxyalkylamino, ω -aminoalkylamino, ω -fosfonoalkylamino, ω -fosfonoalkyloxy, propinyl)-9-isopropylpurine, 2-(R)-(2-hydroxypyrrolidin-1-yl)-6-benzylamino-8-chloro
5 (or hydroxy, bromo, fluoro, amino, amido, carboxy, cyano, methylamino, thio, methylthio, ω -hydroxyalkylamino, ω -hydroxyalkyloxy, ω -carboxyalkylamino, ω -aminoalkylamino, ω -fosfonoalkylamino, ω -fosfonoalkyloxy, propinyl)-9-isopropylpurine, 2-(R)-(1-isopropyl-2-hydroxyethylamino)-6-(3-chloro-4-carboxyanilino)-8-chloro (or hydroxy, bromo, fluoro, amino, amido, carboxy, cyano, methylamino, thio, methylthio,
10 ω -hydroxyalkylamino, ω -hydroxyalkyloxy, ω -carboxyalkylamino, ω -aminoalkylamino, ω -fosfonoalkylamino, ω -fosfonoalkyloxy, propinyl)-9-isopropylpurine, 2-(R)-(1-isopropyl-2-hydroxyethylamino)-6-benzylamino-8-chloro (or hydroxy, bromo, fluoro, amino, amido, carboxy, cyano, methylamino, thio, methylthio, ω -hydroxyalkylamino, ω -hydroxyalkyloxy, ω -carboxyalkylamino, ω -aminoalkylamino, ω -fosfonoalkylamino, ω -fosfonoalkyloxy, propinyl)-9-isopropylpurine, 2-(R)-(1-isopropyl-2-hydroxyethyl-
15 amino)-6-(3-chloroanilino)-8-chloro (or hydroxy, bromo, fluoro, amino, amido, carboxy, cyano, methylamino, thio, methylthio, ω -hydroxyalkylamino, ω -hydroxyalkyloxy, ω -carboxyalkylamino, ω -aminoalkylamino, ω -fosfonoalkylamino, ω -fosfonoalkyloxy, propinyl)-9-isopropylpurine, 2-alkylamino-6-dimethylamino-8-chloro (or hydroxy, bromo, fluoro, amino, amido, carboxy, cyano, methylamino, thio, methylthio, ω -hydroxyalkylamino, ω -hydroxyalkyloxy, ω -carboxyalkylamino, ω -aminoalkylamino, ω -fosfonoalkylamino, ω -fosfonoalkyloxy, propinyl)-9-(R)-(2-phosphonomethoxypropyl)-
20 purine, 2-alkylamino-6-diethylamino-8-chloro (or hydroxy, bromo, fluoro, amino, amido, carboxy, cyano, methylamino, thio, methylthio, ω -hydroxyalkylamino, ω -hydroxyalkyloxy, ω -carboxyalkylamino, ω -aminoalkylamino, ω -fosfonoalkylamino, ω -fosfonoalkyloxy, propinyl)-9-(R)-(2-phosphonomethoxypropyl)purine, 2-alkylamino-6-butylamino-8-chloro (or hydroxy, bromo, fluoro, amino, amido, carboxy, cyano, methylamino, thio, methylthio, ω -hydroxyalkylamino, ω -hydroxyalkyloxy, ω -carboxyalkylamino, ω -aminoalkylamino, ω -fosfonoalkylamino, ω -fosfonoalkyloxy, propinyl)-9-
25 (R)-(2-phosphonomethoxypropyl)purine, 2-alkylamino-6-(2-butylamino)-8-chloro (or hydroxy, bromo, fluoro, amino, amido, carboxy, cyano, methylamino, thio, methylthio, ω -hydroxyalkylamino, ω -hydroxyalkyloxy, ω -carboxyalkylamino, ω -aminoalkylamino, ω -fosfonoalkylamino, ω -fosfonoalkyloxy, propinyl)-9-
30 (R)-(2-phosphonomethoxypropyl)purine, 2-alkylamino-6-(2-butylamino)-8-chloro (or hydroxy, bromo, fluoro, amino, amido, carboxy, cyano, methylamino, thio, methylthio, ω -hydroxyalkylamino, ω -hydroxyalkyloxy, ω -carboxyalkylamino, ω -aminoalkylamino,

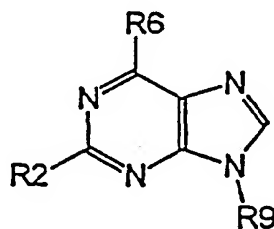
ω -fosfonoalkylamino, ω -fosfonoalkyloxy, propinyl)-9-(R)-(2-phosphonomethoxypropyl)purine, 2-alkylamino-6-cyclopropylamino-8-chloro (or hydroxy, bromo, fluoro, amino, amido, carboxy, cyano, methylamino, thio, methylthio, ω -hydroxyalkylamino, ω -hydroxyalkyloxy, ω -carboxyalkylamino, ω -aminoalkylamino, ω -fosfonoalkylamino, ω -fosfonoalkyloxy, propinyl)-9-(R)-(2-phosphonomethoxypropyl)purine, 2-amino-6-cyclohexylamino-8-chloro (or hydroxy, bromo, fluoro, amino, amido, carboxy, cyano, methylamino, thio, methylthio, ω -hydroxyalkylamino, ω -hydroxyalkyloxy, ω -carboxyalkylamino, ω -aminoalkylamino, ω -fosfonoalkylamino, ω -fosfonoalkyloxy, propinyl)-9-(R)-(2-phosphonomethoxypropyl)purine, 2-alkylamino-6-(pyrrolidin-1-yl)-8-chloro (or hydroxy, bromo, fluoro, amino, amido, carboxy, cyano, methylamino, thio, methylthio, ω -hydroxyalkylamino, ω -hydroxyalkyloxy, ω -carboxyalkylamino, ω -aminoalkylamino, ω -fosfonoalkylamino, ω -fosfonoalkyloxy, propinyl)-9-(R)-(2-phosphonomethoxypropyl)purine, 2-alkylamino-6-(morpholin-1-yl)-8-chloro (or hydroxy, bromo, fluoro, amino, amido, carboxy, cyano, methylamino, thio, methylthio, ω -hydroxyalkylamino, ω -hydroxyalkyloxy, ω -carboxyalkylamino, ω -aminoalkylamino, ω -fosfonoalkylamino, ω -fosfonoalkyloxy, propinyl)-9-(R)-(2-phosphonomethoxypropyl)purine.

26. A method for preparing substituted nitrogen heterocyclic derivatives according to claim 1 of formula I, wherein R2, R6, R8 and R9 are as defined in claim 1, A is a group of formula



25

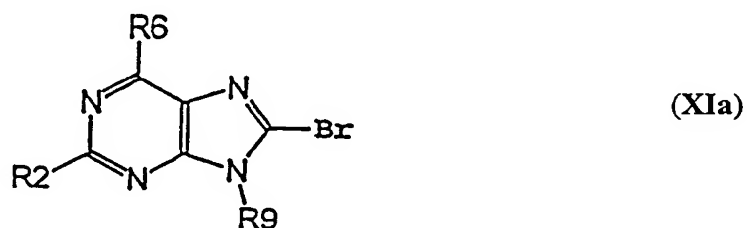
and Z is N, characterized in that a trisubstituted derivative of formula XI,



(XI)

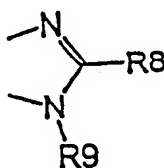
30

wherein R2, R6 and R9 are as defined in claim 1, is brominated with using a brominating system selected from a group consisting of bromine/acetic acid, bromine/chloroform, bromine/acetate buffer, bromine/water, N-bromosuccinimide/dimethylformamide and bromoacetamide/dimethylformamide to obtain a derivative of formula **XIa**,

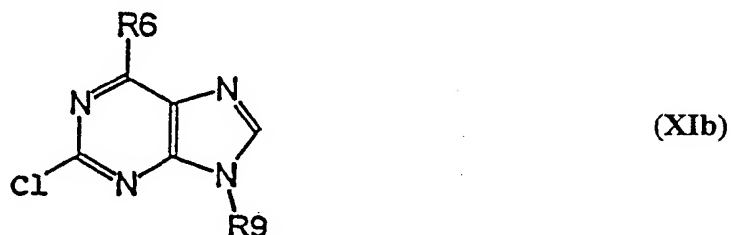


wherein R2, R6 and R9 are as defined in claim 1, and the bromine atom in position 8 of the derivative of formula **XIa** is then optionally subjected to a nucleophilic substitution in order to replace it by another substituent R8 as defined in claim 1.

27. A method for preparing substituted nitrogen heterocyclic derivatives according to claim 1 of formula I, wherein R2, R6, R8 and R9 are as defined in claim 1, A is a group of formula



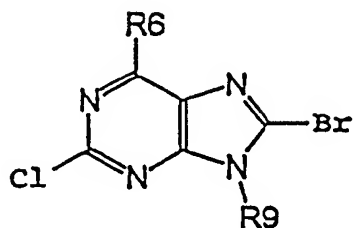
and Z is N, characterized in that a trisubstituted derivative of formula **XIb**,



wherein R6 and R9 are as defined in claim 1, is brominated with using a brominating system selected from a group consisting of bromine/acetic acid, bromine/chloroform, bromine/acetate buffer, bromine/water, N-bromosuccinimide/dimethylformamide and

bromoacetamide/dimethylformamide to obtain a derivative of formula **XIc**,

5

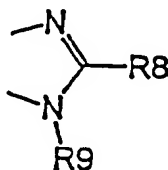


(XIc)

wherein R6 and R9 are as defined in claim 1, and the chlorine atom in position 2 and the bromine atom in position 8 of the derivative of formula **XIc** are optionally, either
10 progressively or simultaneously, subjected to a substitution in order to replace them by another substituents R2 and R8 as defined in claim 1.

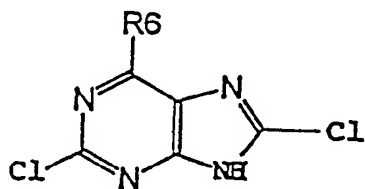
28. A method for preparing substituted nitrogen heterocyclic derivatives of formula I,
wherein R2, R6 and R8 are as defined in claim 1, R9 is alkyl as defined in claim 1, A is
15 a group of formula

20



and Z is N, characterized in that, a trisubstituted derivative of formula **XIIa**,

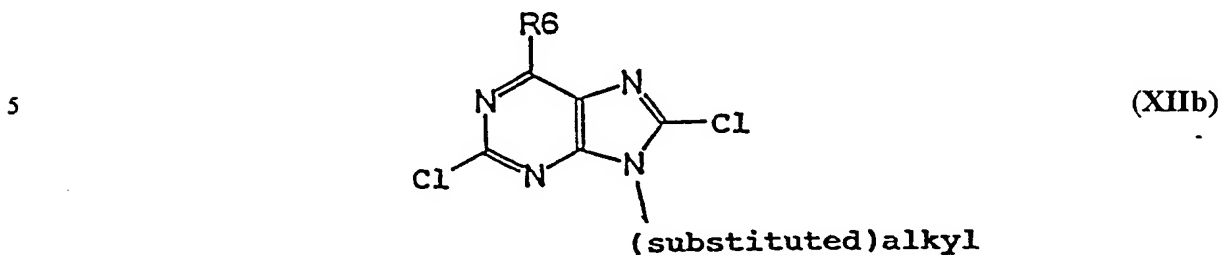
25



(XIIa)

30 wherein R6 is as defined in claim 1, is alkylated in position 9 with using an appropriate alkylating agent in a system selected from a group consisting of K₂CO₃/dimethylformamide, Cs₂CO₃/dimethylformamide, t-BuOK/dimethyl-

formamide, t-BuOK/dimethylsulphoxide and NaH/dimethylformamide or under conditions of Mitsunobu reaction to obtain a derivative of formula **XIIb**,

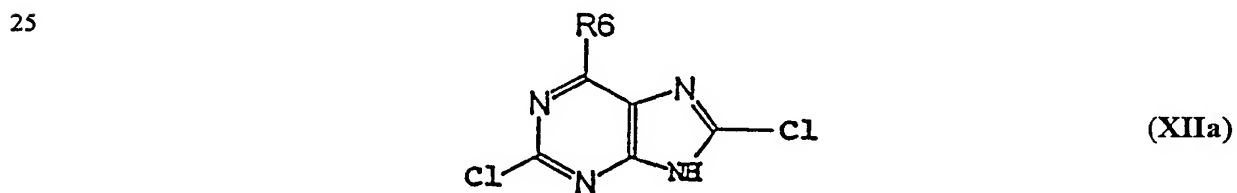


wherein R6 and alkyl are as defined in claim 1, and the chlorine atoms in positions 2 and 8 of the derivative of formula **XIIb** are optionally, either progressively or simultaneously, subjected to a nucleophilic substitution in order to replace them by another substituents R2 and R8 as defined in claim 1.

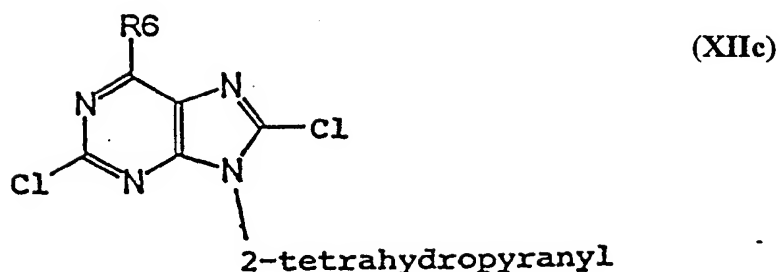
29. A method for preparing substituted nitrogen heterocyclic derivatives according to claim 1, wherein R2, R6 and R8 are as defined in claim 1, R9 is alkyl as defined in claim 1, A is a group of formula



and Z is N, characterized in that a trisubstituted derivative of formula **XIIa**,

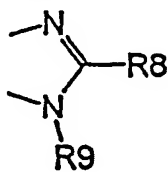


30 wherein R6 is as defined in claim 1, is protected by reacting, for example, with 2-dihydropyranane/ H^+ to obtain a derivative of formula **XIIc**,

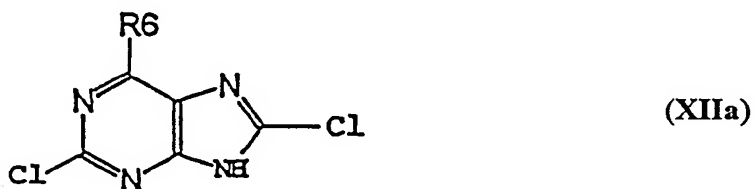


wherein R6 is as defined in claim 1, and a group R6 and/or the chlorine atom in position 2 and/or the chlorine atom in position 8 are optionally converted to another substituents R2, R6 and R8 as defined in claim 1 and in such optionally modified product of formula XIIc, the 2-tetrahydropyranyl group is split off to obtain a corresponding derivative of formula XIIc wherein R9 is hydrogen, and so obtained product is then alkylated in position 9 with using an appropriate alkylating agent in a system selected from a group consisting of K₂CO₃/dimethylformamide, Cs₂CO₃/dimethylformamide, t-BuOK/dimethylformamide, t-BuOK/dimethylsulphoxide and NaH/dimethylformamide or under conditions of Mitsunobu reaction to obtain a tetrasubstituted derivative of formula XIIc, wherein R2, R6, R8 and R9 are as defined in claim 1.

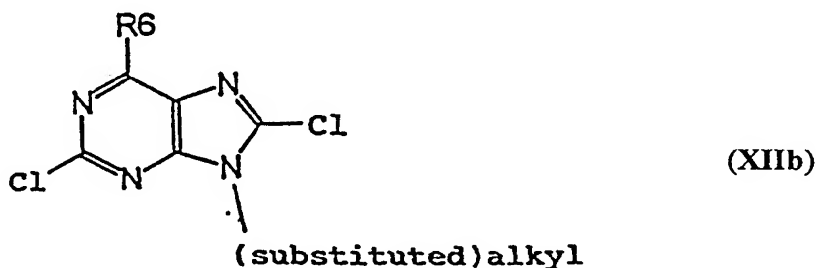
30. A method for preparing substituted nitrogen heterocyclic derivatives according to claim 1 of formula I, wherein R2, R6 and R8 are as defined in claim 1, R9 is alkyl as defined in claim 1, A is a group of formula



and Z is N, characterized in that, 2,6,8-trichloropurine is subjected to a nucleophilic substitution in position 6 in order to replace the chlorine atom in position 6 by another substituent R6 as defined in claim 1 to obtain a derivative of formula XIIa,



wherein R6 is as defined in claim 1, and so obtained product is then alkylated in position 9 with using an appropriate alkylating agent in a system selected from a group consisting of K_2CO_3 /dimethylformamide, Cs_2CO_3 /dimethylformamide, $t\text{-BuOK}$ /dimethylformamide, $t\text{-BuOK}$ /dimethylsulphoxide and NaH /dimethylformamide or under
 5 conditions of Mitsunobu reaction to obtain a derivative of formula XIIb,

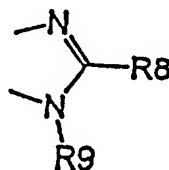


10

wherein R6 and alkyl are as defined in claim 1, and the chlorine atoms in positions 2 and 8 of the derivative of formula XIIb are optionally, either progressively or simultaneously, subjected to a nucleophilic substitution in order to replace them by
 15 another substituents R2 and R8 as defined in claim 1.

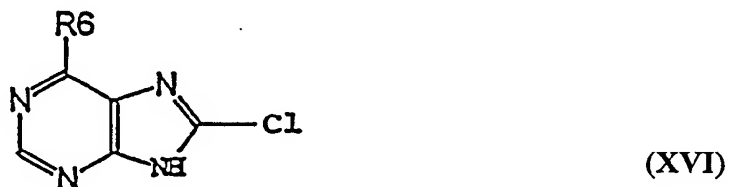
31. A method for preparing substituted nitrogen heterocyclic derivatives according to claim 1 of formula I, wherein R2 is hydrogen, R6 and R8 are as defined in claim 1, R9 is alkyl as defined in claim 1, A is a group of formula

20

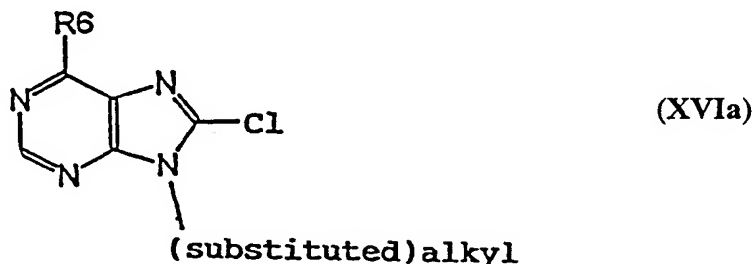


25 and Z is N, characterized in that, 6,8-dichloropurine is subjected to a nucleophilic substitution in position 6 in order to replace the chlorine atom in position 6 by another substituent R6 as defined in claim 1 to obtain a derivative of formula XVI,

30

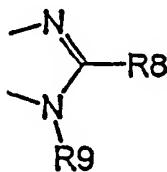


wherein R6 is as defined in claim 1, and so obtained derivative is alkylated in position 9 with using an appropriate alkylating agent in a system selected from a group consisting of K₂CO₃/dimethylformamide, Cs₂CO₃/dimethylformamide, t-BuOK/dimethyl-
 5 formamide, t-BuOK/dimethylsulphoxide and NaH/dimethylformamide or under conditions of Mitsunobu reaction to obtain a derivative of formula XVIa,



wherein R6 and alkyl are as defined in claim 1, and the chlorine atom in position 8 of the derivative of formula XVIa is optionally subjected to a nucleophilic substitution in order
 15 to replace it by another substituent R8 as defined in claim 1.

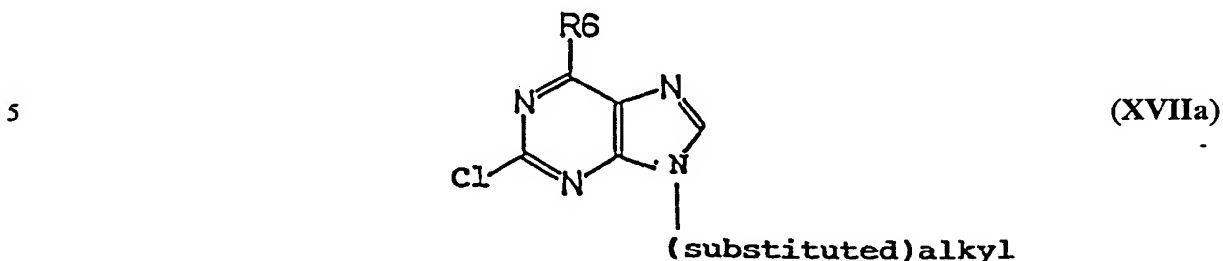
32. A method for preparing substituted nitrogen heterocyclic derivatives according to claim 1 of formula I, wherein R2, R6 and R8 are as defined in claim 1, R9 is alkyl as defined in claim 1, A is a group of formula



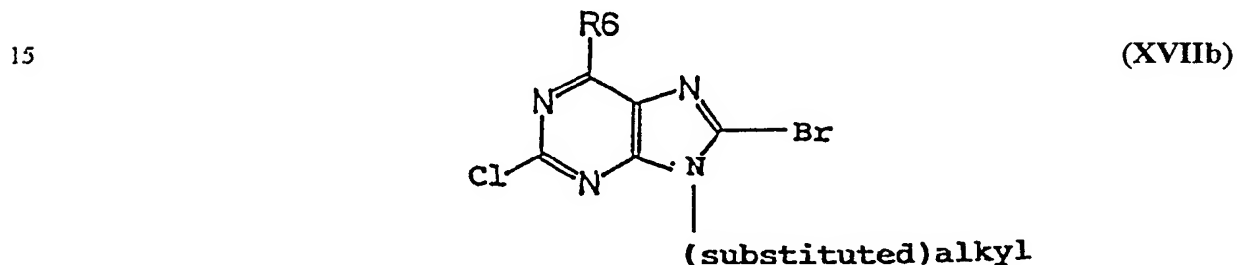
25 and Z is N, characterized in that 2,6-dichloropurine is subjected to a nucleophilic substitution in position 6 in order to replace the chlorine atom in position 6 by another substituent R6 as defined in claim 1, and so obtained derivative of formula XVII,



wherein R6 is as defined in claim 1, is alkylated in position 9 with using either methyl acrylate or acrylonitrile or oxirane to obtain a derivative of formula **XVIIa**,



wherein R6 and alkyl are as defined in claim 1, which product is then brominated in position 8 with using a brominating system selected from a group consisting of bromine/acetic acid, bromine/chloroform, bromine/acetate buffer, bromine/water, N-bromosuccinimide/dimethylformamide and bromoacetamide/dimethylformamide to obtain a derivative of formula **XVIIb**,

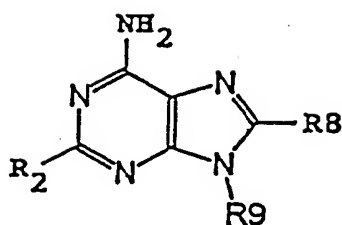


20 wherein R6 and alkyl are as defined in claim 1, and so obtained product is then optionally, either progressively or simultaneously, subjected to a nucleophilic substitution in positions 2 and 8 in order to replace the chlorine atom in position 2 and the bromine atom in position 8 by another substituents R2 and R8 as defined in claim 1.

25 33. A method for preparing substituted nitrogen heterocyclic derivatives according to claim 1 of formula I, wherein R6 is a halogen or hydrogen, R2, R8 and R9 are as defined in claim 1, A is a group of formula



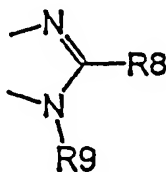
and Z is N, characterized in that a derivative of formula **XVIII**,



(XVIII)

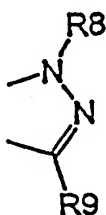
wherein R₂, R₈ and R₉ are as defined in claim 1, is halogenated *via* diazotation with using, for example, amylnitrite/CH₂Br₂ or amylnitrite/CHI₃, and so obtained derivative wherein R₆ is halogen is optionally hydrogenolyzed with using H₂/Pd catalyst to obtain a corresponding derivative wherein R₆ is hydrogen.

34. A method for preparing substituted nitrogen heterocyclic derivatives according to claim 1 of formula I, wherein R₉ is as defined in claim 1 and one substituent of R₂, R₆ or R₈ is alkynyl or alkenyl containing 1 to 6 carbon atoms whereas two other substituents of R₂, R₆ and R₈ are as defined in claim 1, A is a group of



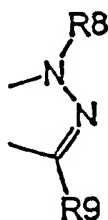
and Z is N or CH, characterized in that a derivative of formula I, wherein R₉ is as defined in claim 1 and one substituent of R₂, R₆ and R₈ is a halogen whereas two other substituents of R₂, R₆ and R₈ are as defined in claim 1, is alkynylated in the position of halogen with using alkine/triphenylphosphine-PdCl₂/CuI/triethylamine system and so obtained 2-, 6- or 8-alkynyl derivative is optionally converted to 2-, 6- or 8-alkenyl derivative with using a Lindlar catalyst.

35. A method for preparing substituted nitrogen heterocyclic derivatives according to claim 1 of formula I, wherein R₂, R₆, R₈ and R₉ are as defined in claim 1, A is a group of formula



and Z is N, characterized in that 2,5-dialkyl-3-alkoxykarbonyl-4-aminopyrazole, wherein alkyls and alkoxy are as defined in claim 1, is reacted with formamidine acetate/triethylamine system to obtain 1,3-dialkyl-7-hydroxypyrazolo[4,3-d]pyrimidine and so obtained 7-hydroxy derivative is then optionally transferred to a corresponding 7-chloro derivative by reaction with thionylchloride, which product is optionally substituted to obtain a corresponding R6-derivative, wherein R6 is as defined in claim 1.

36. A method for preparing substituted nitrogen heterocyclic derivatives according to claim 1 of formula I, wherein R6, R8 and R9 are as defined in claim 1, A is a group of formula



and Z is N, characterized in that 3,7-dialkylpyrazolo[4,3-d]pyrimidine, wherein alkyls are as defined in claim 1, is alkylated with using an appropriate alkylating agent in a system selected from a group consisting of K_2CO_3 /dimethylformamide and Cs_2CO_3 /dimethylformamide to obtain a mixture of trialkylated and tetraalkylated products and R6,R8,R9-trialkylated product is separated by column chromatography.

37. Substituted nitrogen heterocyclic derivatives according to claim 1 of formula I and pharmaceutically acceptable salts thereof for use as medicaments.

38. Substituted nitrogen heterocyclic derivatives according to claim 1 of formula I for use as means for preparing affinity adsorption matrices, immobilized enzymes for process control, immunoassay reagents, diagnostic samples, ^{14}C -, 3H -, avidin- or biotin-labelled compounds, oligonucleotides and diagnostic samples.

39. Substituted nitrogen heterocyclic derivatives according to claim 1 of formula I for use as antimitotic drugs, in particular drugs for elimination or reduction of viral spread or growth in tissue culture systems during the production of biopharmaceutical or

other products such as proteins and vaccines, drugs for elimination or reduction of viral spread and growth in clinical samples such as blood, and drugs for stopping of growth of tissue culture cells while leaving the cells to carry on with protein production.

5 40. A pharmaceutical composition with cytostatic, anticancer, antimitotic, antineurodegenerative, immunosuppressive and antimicrobial activity, characterised in that it comprises, in addition to auxiliary pharmaceutical matters, at least one substituted nitrogen heterocyclic derivative according to claim 1 of formula I.

10 41. A combined pharmaceutical composition with cytostatic effect, characterised in that it comprises, in addition to auxiliary pharmaceutical matters, a combination of at least one substituted nitrogen heterocyclic derivative according to claim 1 of formula I and a cytostatic agent selected from a group consisting of mitoxantron, cis- and carbo-platin, methotrexate, taxol and doxorubicin.

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42. A use of substituted nitrogen heterocyclic derivatives according to claim 1 of formula I for preparing medicaments destined for treating tumours, cancers, psoriasis, rheumatoid arthritis, lupus, type I diabetes, multiple sclerosis, restenosis, polycyclic kidney disease, host graft disease and gout, parasitoses, such as those caused by fungi or
20 prostis, or Alzheimer's disease, or for preparing antineurodegenerative drugs and suppress immunostimulation agents.

INTERNATIONAL SEARCH REPORT

International Application No

PCT/CZ 00/00002

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C07D473/16 C07D473/32 C07D473/34 C07D473/40 C07D487/04
A61K31/52 A61P35/00 //(C07D487/04, 239:00, 231:00),
(C07D487/04, 239:00, 209:00)

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C07D C07F A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 98 01448 A (JAPAN ENERGY CORP ;CHIBA NOBUYOSHI (JP); HIROTA KOHSAKU (JP); ISOB) 15 January 1998 (1998-01-15) cited in the application claims	1-42
Y	CA 2 231 005 A (INSTITUTE OF EXPERIMENTAL BOTANY OF THE ACADEMY OF SCIENCES OF THE CZE) 4 September 1999 (1999-09-04) claims	1-42
Y	WO 97 20842 A (CENTRE NAT RECH SCIENT ;MEIJER LAURENT (FR); BISAGNI EMILE (FR); L) 12 June 1997 (1997-06-12) cited in the application claims	1-42
	-/--	

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance

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"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

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Date of the actual completion of the international search

12 May 2000

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INTERNATIONAL SEARCH REPORT

International Application No

PCT/CZ 00/00002

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	WO 97 16452 A (CIBA GEIGY AG ;ZIMMERMANN JUERG (CH); CAPRARO HANS GEORG (CH); PET) 9 May 1997 (1997-05-09) cited in the application claims -----	1-42
Y	WO 98 05335 A (CV THERAPEUTICS INC ;BLUM CHERI LYNN (US); LUM ROBERT T (US); MACK) 12 February 1998 (1998-02-12) cited in the application claims -----	1-42
Y	WO 96 33200 A (USTAV ORGANICKE CHEMIE A BIOCH ;REGA STICHTING (BE); HOLY ANTONIN) 24 October 1996 (1996-10-24) cited in the application claims -----	1-42
P,X	WO 99 28321 A (SUMITOMO PHARMACEUTICALS CO.,LTD.) 10 June 1999 (1999-06-10) claims -----	1-42
P,Y	WO 99 07705 A (MEIJER LAURENT ;SCHULTZ PETER (US); KIM SUNG HOU (US); UNIV CALIFO) 18 February 1999 (1999-02-18) claims -----	1-42

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/CZ 00/00002

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
WO 9801448	A	15-01-1998	AU 698419 B	29-10-1998
			AU 3358497 A	02-02-1998
			CA 2230808 A	15-01-1998
			EP 0882727 A	09-12-1998
			NO 980908 A	30-04-1998
			NZ 329798 A	29-04-1999
			US 6028076 A	22-02-2000
CA 2231005	A		NONE	
WO 9720842	A	12-06-1997	FR 2741881 A	06-06-1997
			CA 2238843 A	12-06-1997
			EP 0874847 A	04-11-1998
			JP 2000501408 T	08-02-2000
WO 9716452	A	09-05-1997	AU 7296896 A	22-05-1997
			BR 9611157 A	30-03-1999
			CA 2234609 A	09-05-1997
			CN 1202896 A	23-12-1998
			EP 0874846 A	04-11-1998
			JP 11514336 T	07-12-1999
WO 9805335	A	12-02-1998	US 5866702 A	02-02-1999
			AU 3900097 A	25-02-1998
			CN 1231611 A	13-10-1999
			NO 990466 A	25-03-1999
			PL 331408 A	19-07-1999
WO 9633200	A	24-10-1996	US 5977061 A	02-11-1999
			AU 5268696 A	07-11-1996
			EP 0821690 A	04-02-1998
			JP 11511114 T	28-09-1999
WO 9928321	A	10-06-1999	AU 1260299 A	16-06-1999
WO 9907705	A	18-02-1999	AU 8773098 A	01-03-1999